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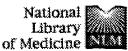
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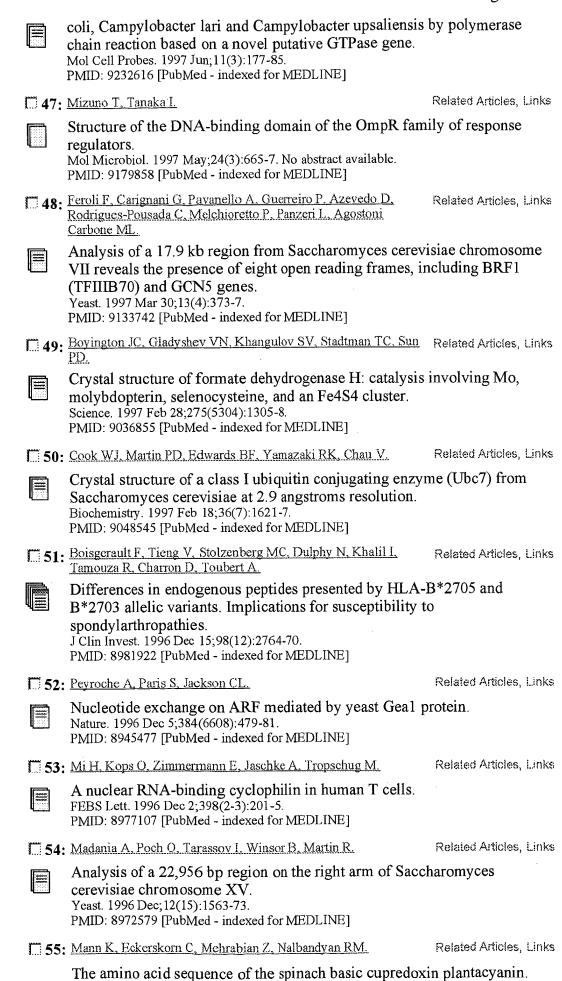
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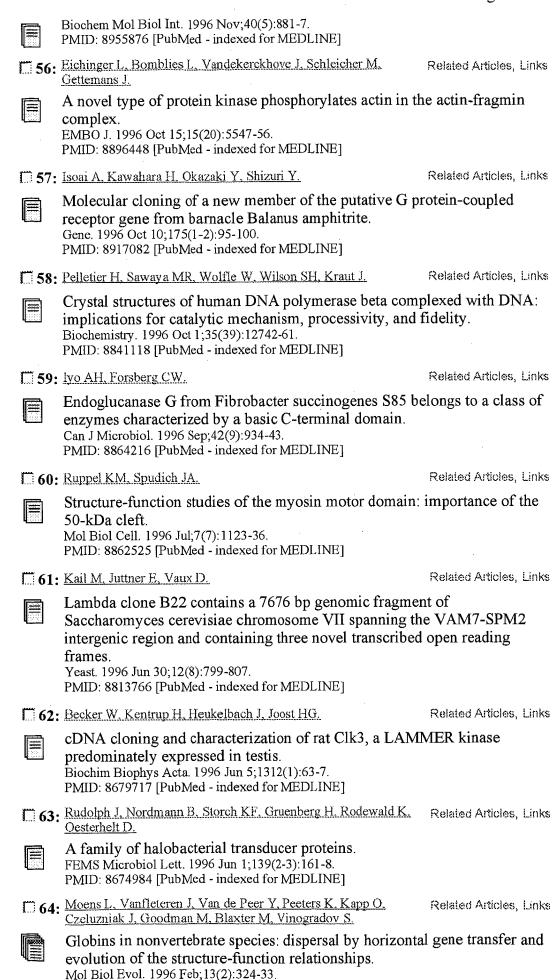
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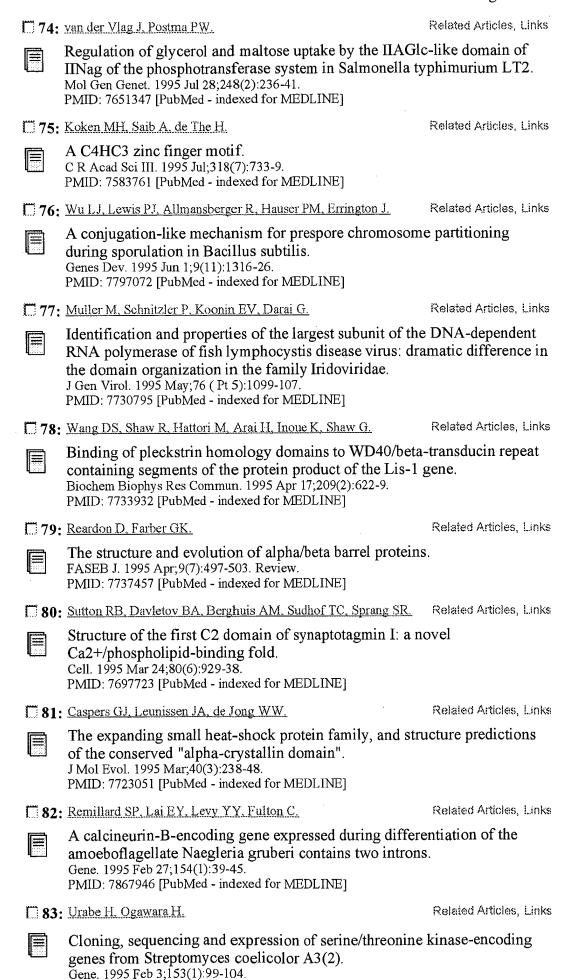
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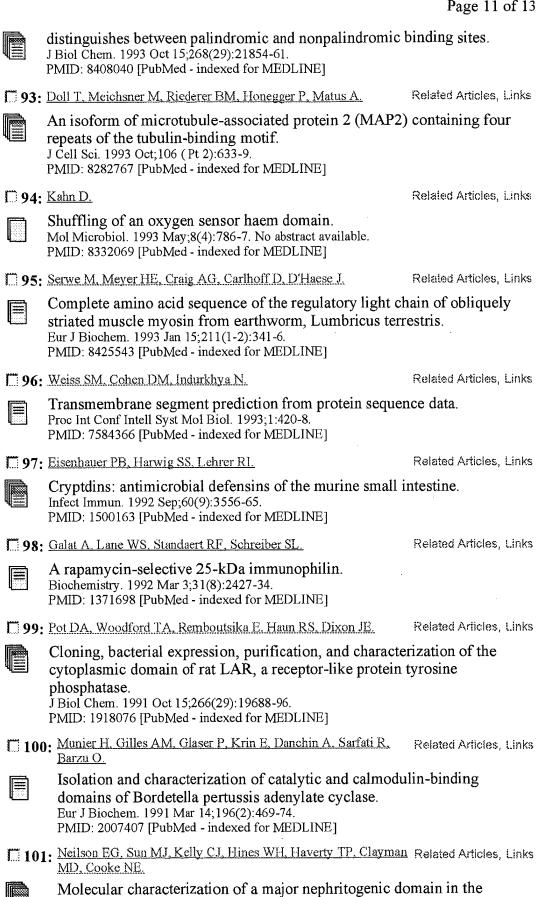
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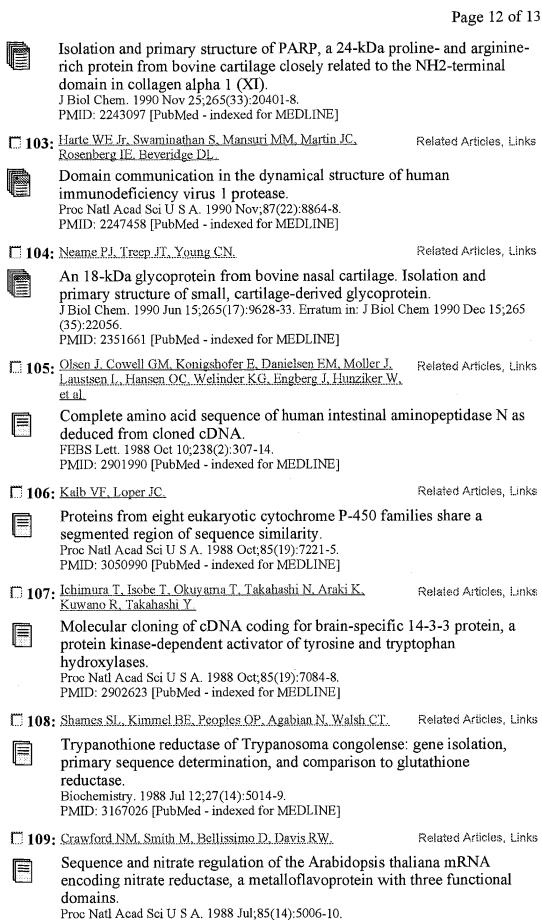


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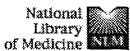
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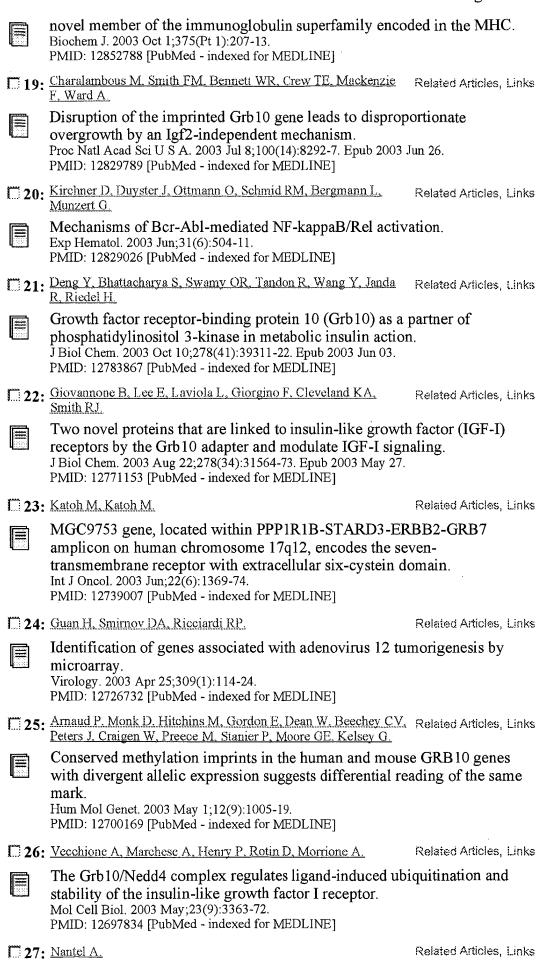
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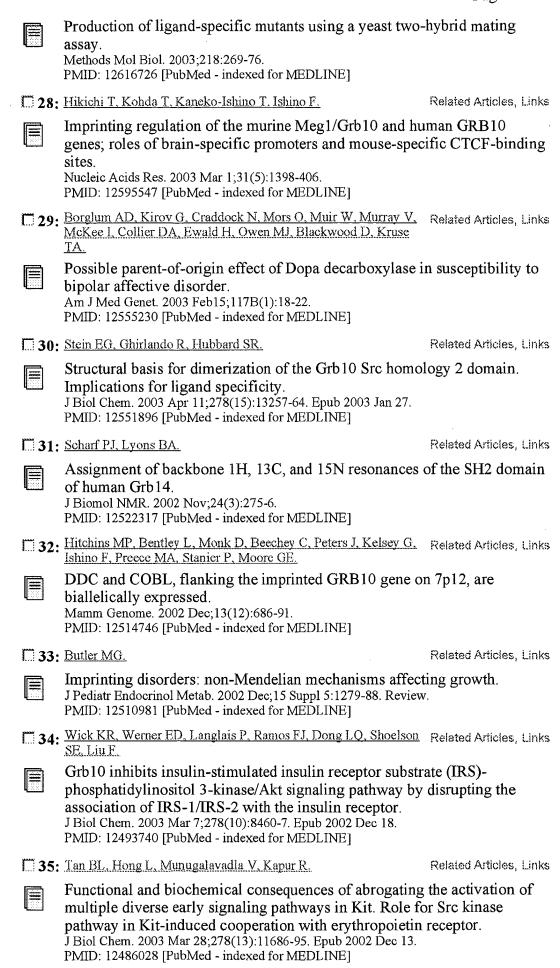


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     139:287335
     Gene expression profiling in the brain of rat models and use of nucleotide
TI
     sequences as gene chips for screening antidepressants
     Yoshikawa, Takeo; Nakaya, Noriaki; Aburaya, Hiroyuki
Institute of Physical and Chemical Research, Japan
IN
PA
     Jpn. Kokai Tokkyo Koho, 18 pp.
SO
     CODEN: JKXXAF
DT
     Patent
     Japanese
LA
FAN.CNT 1
     PATENT NO.
                      KIND DATE
                                            APPLICATION NO.
                                                              DATE
PΙ
     JP 2003274949
                       A2
                             20030930
                                            JP 2002-81502
                                                              20020322
PRAI JP 2002-81502
                             20020322
L6
     ANSWER 7 OF 135 IFIPAT COPYRIGHT 2004 IFI on STN DUPLICATE 2
ΑN
      10399185 IFIPAT; IFIUDB; IFICDB
      DIAGNOSIS OF DISEASES ASSOCIATED WITH THE IMMUNE SYSTEM BY DETERMINING
TI
      CYTOSINE METHYLATION
IN
      Berlin Kurt (DE); Olek Alexander (DE); Piepenbrock Christian (DE)
      Unassigned Or Assigned To Individual (68000)
PA
                      A1 20030731
PΙ
      us 2003143606
      US 2002-311455
ΑI
                           20021216
      WO 2001-EP7537
                           20010702
                           20021216
                                     PCT 371 date
                           20021216
                                     PCT 102(e) date
      DE 2000-10032529
PRAI
                           20000630
      DE 2000-10043826
                           20000901
FI
      US 2003143606
                           20030731
DT
      Utility; Patent Application - First Publication
FS
      CHEMICAL
      APPLICATION
CLMN
      31
L6
     ANSWER 8 OF 135 IFIPAT COPYRIGHT 2004 IFI ON STN DUPLICATE 3
      10385219
ΑN
                IFIPAT; IFIUDB; IFICDB
TI
      GDU, A NOVEL SIGNALLING PROTEIN
IN
      Daly Roger John (AU); Sutherland Robert Lyndsay (AU)
      Unassigned Or Assigned To Individual (68000)
PA
PΙ
      US 2003129639
                      A1 20030710
ΑI
      us 2002-323001
                           20021218
                           19960502 Section 371 PCT Filing UNKNOWN
RLI
      WO 1996-AU258
      US 1998-945771
                           19980422 DIVISION
                                                            6465623
      US 2002-242332
                           20020911 DIVISION
PRAI
      AU 1995-2742
                           19950502
FI
      US 2003129639
                           20030710
      US 6465623
DT
      Utility; Patent Application - First Publication
      CHEMICAL
FS
      APPLICATION
CLMN
GI
       3 Figure(s).
     FIG. 1 shows a schematic representation of ***Grb14***
                                                                  structure with
      a restriction map for the ***Grb14*** cDNA and the cDNA clones used
                      ***Grb14***
                                     sequence aligned underneath. The initial
      to derive the
      clone isolated by CORT screening was designated clone 1. Two other clones
      (1-1 and 1-2) were isolated from the 184 cell line library by screening
                                       ***Grb14***
      using clone 1 as a probe. The
                                                     CDNA sequence was completed
      using two clones L5 and L6, isolated from a human liver cDNA library.
      Abbreviations are as follows: A; Apa I; Av; Avr II, X; Xho I; E; Eco RI.
      The numbers refer to distance in bp.
     FIG. 2 shows the nucleotide and amino acid sequence of
                                                                ***Grb14***
      The PH domain is underlined and the SH2 domain indicated by bold type.
      The translation termination codon is shown by an asterisk in the amino
      acid sequence. Numbers refer to distances in bp.
     FIG. 3 shows the sequence homology between
                                                  ***Grb14***
```

, Grb7, Grb10

and F10E9.6. As alignment of the amino acid sequences of ***Grb14***, mouse Grb7, mouse Grb10 and C. elegans F10E9.6 was obtained using the computer programs Clustal W and SeqVu. Identical residues are boxed. A highly conserved proline-rich motif is indicated by the dotted underline, the PH domain by the broken underline and the SH2 domain by the bold underline. Only the central region of F10E9.6 exhibiting homology with the Grb7 family is shown. Amino acid residues for each protein are numbered (from the initiation methionine) on the right.

ANSWER 9 OF 135 IFIPAT COPYRIGHT 2004 IFI on STN DUPLICATE 4 L6 10360583 IFIPAT; IFIUDB; IFICDB AN METHODS AND COMPOSITIONS FOR INHIBITING GRB7 TI Krag David N; Oligino Lyn; Pero Stephanie C IN Unassigned Or Assigned To Individual (68000) PA A1 20030605 us 2003105000 ΡI us 2001-13815 20011105 ΑI US 2000-245755P 20001103 (Provisional) PRAI us 2003105000 20030605 FI Utility; Patent Application - First Publication DT CHEMICAL FS **APPLICATION** CLMN 93 GI 16 Figure(s). FIG. 1 is a histogram showing the binding of Grb7 binding peptides (G7BP) to the SH2 domain of human Grb7 by ELISA. FIG. 2 is a histogram showing the binding of a control phage clone to the SH2 domain of Grb2 but not to the SH2 domain of Grb7. FIG. 3 is a histogram showing the effect of mutation on a G7BP-4 phage clone on its ability to bind to the SH2 domain of Grb7. FIG. 4 is a histogram showing the binding specificity of seven Grb7 binding peptides to the SH2 domains of Grb7, Grb7 beta D5beta D6, ***Grb14*** , full length Grb2, and BSA using a phage ELISA. FIG. 5A is a graph showing the inhibition of G7-18 peptide-phage binding to Grb7-SH2 with the free synthetic peptides G7-18, G718NATE and G7-8. FIG. 5B is a graph showing the inhibition of G7-8NA peptidephage binding to Grb7-SH2 with the free synthetic peptides G7-8, G7-8NA and G7-8NATE. FIG. 6A is a graph showing G7-18NATE inhibits the association of Grb7 with the ErbB family of receptors, as detected by antiphosphotyrosine. FIG. 6B is a densitometric analysis of autoradiographs using the Biorad Fluor-S Multimager with Quantity One 4.2.1 software, showing G7-18NATE inhibits the association of Grb7, not Grb2, with the ErbB family in a dose-dependent manner. FIG. 7A is a graph showing that G7-18NATE inhibits the association of Grb7 with ErbB3 specifically in a dose-dependent manner, as detected by anti-ErbB3 FIG. 7B is a densitometric analysis of autoradiographs using the Biorad Fluor-S Multimager with Quantity One 4.2.1 software showing that G7-18NATE inhibits the association of Grb7 with ErbB3 in a dose-dependent manner. FIG. 8A is a graph showing that G7-18NATE inhibits the association of Grb7 with ErbB2 specifically in a dose-dependent manner, as detected by anti-ErbB2. FIG. 8B is a densitometric analysis of autoradiographs using the Biorad Fluor-S Multimager with Quantity One 4.2.1 software showing that G7-18NATE inhibits the association of Grb7 with ErbB2 in a dose-dependent FIG. 9A is one possible chemical structure for G7BP-4NATE (SEQ ID NO:50). Other thioether linkages are illustrated in FIGS. 9B, 9C, 9D and 9E, and it is to be understood that any of these linkages can be used in the formation of G7BP-4NATE. FIG. 9B is the structure of a thioether containing peptide (G1TE) . This structure illustrates one possible thioether linkage between the N and C terminals of a peptide that can be used in the thioether containing peptides of the invention. FIG. 9C is another possible structure for the thioether containing peptide GI TE. This structure illustrates one possible thioether linkage between the N and C terminals of a peptide that can be used in the thioether containing peptides of the invention. FIG. 9D is another possible structure for the thioether containing peptide G1TE. This structure illustrates one possible thioether linkage between the N and C terminals of a peptide that can be used in the thioether containing peptides of the invention. FIG. 9E is another possible structure for the thioether containing peptide GI TE. This structure illustrates one possible thioether linkage between the N and C terminals of a peptide that can be used in the thioether

containing peptides of the invention.

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L6
       ANSWER 10 OF 135 IFIPAT COPYRIGHT 2004 IFI ON STN DUPLICATE 5
        10300425 IFIPAT; IFIUDB; IFICDB
ΑN
        GDU, A NOVEL SIGNALLING PROTEIN
TI
        Daly Roger John (AU); Sutherland Robert Lyndsay (AU)
IN
        Unassigned Or Assigned To Individual (68000)
PA
                              A1 20030306
ΡI
        US 2003044834
                                    20020911
ΑI
        US 2002-242332
        WO 1996-AU258
                                    19960502 Section 371 PCT Filing UNKNOWN
RLI
        US 1998-945771
                                    19980422 DIVISION
        AU 1995-2742
                                    19950502
PRAI
                                    20030306
FΙ
        us 2003044834
        us 6465623
        Utility; Patent Application - First Publication
DT
        CHEMICAL
FS
        APPLICATION
CLMN
          3 Figure(s).
GΙ
       FIG. 1 shows a schematic representation of ***Grb14***
                                                                                         structure with
        a restriction map for the ***Grb14*** cDNA and the cDNA clones use to derive the ***Grb14*** sequence aligned underneath. The initial
                                                                   cDNA and the cDNA clones used
        clone isolated by CORT screening was designated clone 1. Two other clones (1-1 and 1-2) were isolated from the 184 cell line library by screening using clone 1 as a probe. The ***Grb14*** cDNA sequence was completed using two clones L5 and L6, isolated from a human liver cDNA library.
        Abbreviations are as follows: A; Apa I; AV; AVr II, X; Xho I; E; Eco RI.
        The numbers refer to distance in bp.
                                                                                       ***Grb14***
       FIG. 2 shows the nucleotide and amino acid sequence of
        The PH domain is underlined and the SH2 domain indicated by bold type.
        The translation termination codon is shown by an asterisk in the amino
        acid sequence. Numbers refer to distances in bp.
        and F10E9.6. As alignment of the amino acid sequences of ***Grb14***, mouse Grb7, mouse Grb10 and C. elegans F10E9.6 was obtained using the computer programs Clustal W and SeqVu. Identical residues are boxed. A highly conserved proline-rich motif is indicated by the dotted underline, the PH domain by the broken underline and the SH2 domain by the bold underline. Only the central region of F10F9 6 exhibiting boxelegy with
       FIG. 3 shows the sequence homology between
        underline. Only the central region of F10E9.6 exhibiting homology with
        the Grb7 family is shown. Amino acid residues for each protein are
        numbered (from the initiation methionine) on the right.
       ANSWER 11 OF 135 USPATFULL ON STN
L6
          2003:330145 USPATFULL
ΑN
          Skin cell biomarkers and methods for identifying biomarkers using
ΤI
          nucleic acid microarrays
         Dooley, Thomas P., Vestavia Hills, AL, UNITED STATES Curto, Ernest V., Huntsville, AL, UNITED STATES Davis, Richard L., JR., Homewood, AL, UNITED STATES
IN
          US 2003232356
PΙ
                                            20031218
                                    Α1
          US 2003-361006
                                            20030210 (10)
ΑI
                                    Α1
PRAI
          US 2002-354519P
                                     20020208 (60)
          Utility
DT
FS
          APPLICATION
LN.CNT 1897
          INCLM: 435/006.000
INCL
          INCLS: 702/020.000
                   435/006.000
NCL
          NCLM:
          NCLS:
                   702/020.000
IC
          [7]
          ICM: C12Q001-68
          ICS: G06F019-00; G01N033-48; G01N033-50
       ANSWER 12 OF 135 USPATFULL on STN
L6
AN
          2003:225702 USPATFULL
          Polynucleotide encoding a novel pleckstrin homology domain and proline
TI
          rich domain containing adapter protein, PMN29
          Finger, Joshua N., San Marcos, CA, UNITED STATES
Perez-Villar, Juan J., Mercerville, NJ, UNITED STATES
IN
         Rajashekar, Reddy, Langhorne, PA, UNITED STATES Yang, Guchen, Morrisville, PA, UNITED STATES Kiener, Peter A., Doylestown, PA, UNITED STATES US 2003157514 A1 20030821
PΙ
          us 2002-234816
                                            20020904 (10)
ΑI
                                    Α1
          US 2001-317063P
                                     20010904 (60)
PRAI
DT
         Utility
          APPLICATION
LN.CNT 13865
```

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INCL
       INCLM: 435/006.000
       INCLS: 435/069.100; 435/320.100; 435/325.000; 530/350.000; 536/023.500;
              435/007.200
NCL
       NCLM:
              435/006.000
              435/069.100; 435/320.100; 435/325.000; 530/350.000; 536/023.500;
       NCLS:
              435/007.200
       [7]
IC
       ICM: C12Q001-68
       ICS: G01N033-53; G01N033-567; C07H021-04; C12P021-02; C12N005-06;
       C07K014~47
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 13 OF 135 USPATFULL ON STN
L6
       2003:37578 USPATFULL
ΑN
       Specimen-linked G protein coupled receptor database
TI
       Muraca, Patrick J., Pittsfield, MA, UNITED STATES
IN
ΡI
       US 2003027223
                          Α1
                                20030206
       US 2002-184694
                                20020628 (10)
                          Α1
ΑI
PRAI
       US 2001-302316P
                           20010629 (60)
       Utility
DT
       APPLICATION
FS
LN.CNT 3618
       INCLM: 435/007.210
INCL
       INCLS: 702/019.000
              435/007.210
NCL
       NCLM:
              702/019.000
       NCLS:
IC
       [7]
       ICM: G01N033-567
       ICS: G06F019-00; G01N033-48; G01N033-50
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 14 OF 135 USPATFULL ON STN
L6
       2003:222015
                    USPATFULL
AN
       Compositions for the detection of blood cell and immunological response
TI
       gene expression
       Cocks, Benjamin G., Sunnyvale, CA, United States
IN
       Stuart, Susan G., Montara, CA, United States
       Seilhamer, Jeffrey J., Los Altos Hills, CA, United States
       Incyte Corporation, Palo Alto, CA, United States (U.S. corporation)
PA
                          в1
PΙ
       US 6607879
                                20030819
                                19980209 (9)
       us 1998-23655
ΑI
       Utility
DT
FS
       GRANTED
LN.CNT
       3719
       INCLM: 435/006.000
INCL
       INCLS: 435/069.100; 536/023.100; 536/024.100; 536/024.300; 536/024.310;
              536/024.320; 536/024.330
              435/006.000
NCL
       NCLM:
              435/069.100; 536/023.100; 536/024.100; 536/024.300; 536/024.310;
       NCLS:
              536/024.320; 536/024.330
       [7]
IC
       ICM: C12Q001-68
       ICS: C07H021-00
       435/6; 435/69.1; 536/22.1; 536/23.1; 536/24.1; 536/24.3-24.33
FXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 15 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN
L6
     DUPLICATE 6
     2003:254017
ΑN
                  BIOSIS
     PREV200300254017
DN
TI
     Structural basis for dimerization of the Grb10 Src homology 2 domain.
     Implications for ligand specificity.
     Stein, Evan G.; Ghirlando, Rodolfo; Hubbard, Stevan R. [Reprint Author]
ΑU
     Skirball Institute of Biomolecular Medicine, New York University School of
CS
     Medicine, 540 First Ave., New York, NY, 10016, USA
     hubbard@saturn.med.nyu.edu
50
     Journal of Biological Chemistry, (April 11, 2003) Vol. 278, No. 15, pp.
     13257-13264. print.
     CODEN: JBCHA3. ISSN: 0021-9258.
     Article
DT
     English
LA
ED
     Entered STN: 28 May 2003
     Last Updated on STN: 28 May 2003
     ANSWER 16 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. ON STN
L6
     DUPLICATE 7
```

```
ΑN
     2003:355616 BIOSIS
     PREV200300355616
DN
     NIK is a component of the EGF/heregulin receptor signaling complexes.
TI
     Chen, Danying; Xu, Liang-Guo; Chen, Lei; Li, Lixia; Zhai, Zhonghe; Shu,
ΑU
     Hong-Bing [Reprint Author]
     Department of Immunology, National Jewish Medical and Research Center, University of Colorado Health Sciences Center, 1400 Jackson Street, K516c,
CS
      Denver, CO, 80206, USA
      shuh@njc.org
     Oncogene, (10 July, 2003) Vol. 22, No. 28, pp. 4348-4355. print.
SO
     ISSN: 0950-9232 (ISSN print).
     Article
DT
     English
LA
     Entered STN: 6 Aug 2003
ED
      Last Updated on STN: 6 Aug 2003
     ANSWER 17 OF 135 LIFESCI
                                         COPYRIGHT 2004 CSA on STN
L6
      2003:53872 LIFESCI
ΑN
     The Grb10/Nedd4 Complex Regulates Ligand-Induced Ubiquitination and
TI
      Stability of the Insulin-Like Growth Factor I Receptor
     Vecchione, A.; Marchese, A.; Henry, P.; Rotin, D.; Morrione, A.*
Department of Urology and Kimmel Cancer Center, Thomas Jefferson
CS
     University, BLSB Room 631, 233 South 10th St., Philadelphia, PA 19107; E-mail: Andrea.Morrione@mail.tju.edu
     Molecular and Cellular Biology [Mol. Cell. Biol.], (20030500) vol. 23, no.
S<sub>0</sub>
      9, pp. 3363-3372.
      ISSN: 0270-7306.
DT
      Journal
FS
LA
      English
SL
      English
L6
      ANSWER 18 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN
      2003:476188 BIOSIS
ΑN
      PREV200300476188
DN
TI
      Characterization of a novel gene (HGP1) potentially involved in
      osteosarcoma progression.
      Eppert, Kolja [Reprint Author]; Aneliunas, Vicky [Reprint Author]; Wunder,
ΑU
      Jay S. [Reprint Author]; Andrulis, Irene L. [Reprint Author]
      Fred A. Litwin Centre for Cancer Genetics, Samuel Lunenfeld Research
CS
     Institute, Mount Sinai Hospital, Toronto, ON, Canada
Proceedings of the American Association for Cancer Research Annual
Meeting, (July 2003) Vol. 44, pp. 1041-1042. print.
Meeting Info.: 94th Annual Meeting of the American Association for Cancer
S0
      Research. Washington, DC, USA. July 11-14, 2003.
      ISSN: 0197-016X.
      Conference; (Meeting)
Conference; Abstract; (Meeting Abstract)
DT
LA
      English
      Entered STN: 15 Oct 2003
ED
      Last Updated on STN: 15 Oct 2003
       ANSWER 19 OF 135 BIOTECHNO COPYRIGHT 2004 Elsevier Science B.V. on STN
L6
       DUPLICATE
       2003:37386423
                           BIOTECHNO
ΑN
       Using gene expression profiling to identify the molecular basis of the synergistic actions of hepatocyte growth factor and vascular endothelial
TI
       growth factor in human endothelial cells
       Gerritsen M.E.; Tomlinson J.E.; Zlot C.; Ziman M.; Hwang S.
M.E. Gerritsen, 541 Parrott Drive, San Mateo, CA 94402, United States.
ΑU
CS
       E-mail: meg570@comcast.net
       British Journal of Pharmacology, (2003), 140/4 (595-610), 52 reference(s)
S0
       CODEN: BJPCBM ISSN: 0007-1188
       Journal; Conference Article
DT
CY
       United Kingdom
       English
LA
SL
       English
      ANSWER 20 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN
L6
      DUPLICATE 9
      2003:451765
AN
                     BIOSIS
      PREV200300451765
DN
TI
      Carcinogen mediated initiation of glial progenitors in the rat brain
      results in marked dependency of proliferation and differentiation by
      insulin and FGF-2.
      Kokkinakis, Demetrius Michael [Reprint Author]; Yang, Shuting [Reprint
ΑU
```

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Author]
     University of Pittsburgh, Pittsburgh, PA, USA
CS
     Proceedings of the American Association for Cancer Research Annual
S0
     Meeting, (July 2003) Vol. 44, pp. 482. print.
Meeting Info.: 94th Annual Meeting of the American Association for Cancer
     Research. Washington, DC, USA. July 11-14, 2003.
     ISSN: 0197-016x.
     Conference; (Meeting)
DT
     Conference; Abstract; (Meeting Abstract)
     English
LA
ED
     Entered STN: 1 Oct 2003
     Last Updated on STN: 1 Oct 2003
     ANSWER 21 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. ON STN
L6
     DUPLICATE 10
     2003:587917
                   BIOSIS
AΝ
     PREV200300570714
DN
                                          is an intrinsically unstructured
                           ***Grb14***
TI
     The PIR domain of
     protein: Implication in insulin signaling.
     Moncoq, Karine; Broutin, Isabelle [Reprint Author]; Larue, Valery;
ΑIJ
     Perdereau, Dominique; Cailliau, Katia; Browaeys-Poly, Edith; Burnol,
     Anne-Francoise; Ducruix, Arnaud
Laboratoire de Cristallographie et RMN Biologiques, Faculte de Pharmacie
Paris 5, 4 avenue de l'Observatoire, 75270, Paris Cedex, 06, France
CS
     broutin@pharmacie.univ-paris5.fr
     FEBS Letters, (20 November 2003) Vol. 554, No. 3, pp. 240-246. print.
S0
     CODEN: FEBLAL. ISSN: 0014-5793.
     Article
DT
     English
LA
     Entered STN: 10 Dec 2003
ED
     Last Updated on STN: 10 Dec 2003
     ANSWER 22 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. ON STN
L6
     DUPLICATE 11
     2003:390017
                   BIOSIS
ΑN
     PREV200300390017
DN
     Inhibition of FGF receptor signalling in Xenopus oocytes: Differential
TI
     effect of Grb7, Grb10 and ***Grb14***
     Cailliau, Katia; Le Marcis, Veronique; Bereziat, Veronique; Perdereau,
ΑU
     Dominique: Cariou, Bertrand: Vilain, Jean Pierre; Burnol, Anne-Francoise;
     Browaeys-Poly, Edith [Reprint Author]
     Laboratoire de Biologie du Developpement, Universite des Sciences;
CS
     Technologies de Lille, UPRES UA 1033, IFR 118, Batiment SN3, Villeneuve
     d'Ascq Cedex, France
     edith.browaeys@univ-lille1.fr
     FEBS Letters, (31 July, 2003) Vol. 548, No. 1-3, pp. 43-48. print. CODEN: FEBLAL. ISSN: 0014-5793.
S0
DT
     Article
     English
LA
ED
     Entered STN: 27 Aug 2003
     Last Updated on STN: 27 Aug 2003
     ANSWER 23 OF 135 IFIPAT COPYRIGHT 2004 IFI on STN DUPLICATE 12
L6
ΑN
       10142689 IFIPAT; IFIUDB; IFICDB
       GDU, A NOVEL SIGNALLING PROTEIN; IT MAY PROVIDE A TARGET IN DISEASES OR
TI
       CONDITIONS IN WHICH PLATELET DERIVED GROWTH FACTOR RECEPTOR (PDGFR) PLAYS
       A REGULATORY ROLE E.G. WOUND HEALING, FIBROTIC CONDITIONS,
       ATHEROSCLEROSIS
       DALY ROGER JOHN (AU); SUTHERLAND ROBERT LINDSAY (AU)
IN
       Unassigned Or Assigned To Individual (68000)
PA
PPA
       Garvan Institute of Medical Research AU (Probable)
PΙ
       US 2002086328
                            20020704
                        Α1
       US 1998-945771
                             19980422
ΑI
       wo 1996-us258
                             19960502
       US 2002086328
                             20020704
FI
                             20021015
       us 6465623
       Utility; Patent Application - First Publication
DT
FS
       CHEMICAL
       APPLICATION
CLMN
        3 Figure(s).
GI
                                                       ***Grb14***
      FIG. 1 shows a schematic representation of
                                                                      structure with
       a restriction map for the ***Grb14*** to derive the ***Grb14*** sequence
                                                     cDNA and the cDNA clones used
                                      sequence aligned underneath. The initial
       clone isolated by CORT screening was designated clone 1. Two other clones
```

(1-1 and 1-2) were isolated from the 184 cell line library by screening

```
Abbreviations are as follows: A; Apa I; Av; Avr II, X; Xho I; E; Eco RI.
       The numbers refer to distance in bp.
      FIG. 2 shows the nucleotide and amino acid sequence of
                                                                           ***Grb14***
       The PH domain is underlined and the SH2 domain indicated by bold type.
       The translation termination codon is shown by an asterisk in the amino
      acid sequence. Numbers refer to distances in bp. FIG. 3 shows the sequence homology between ***G
       and F10E9.6. As alignment of the amino acid sequences of ***Grb14***
                                                            ***Grb14***
       mouse Grb7, mouse Grb10 and C. elegans F10E9.6 was obtained using the
       computer programs Clustal W and SeqVu. Identical residues are boxed. A
       highly conserved proline-rich motif is indicated by the dotted underline,
       the PH domain by the broken underline and the SH2 domain by the bold
       underline. Only the central region of F10E9.6 exhibiting homology with
       the Grb7 family is shown. Amino acid residues for each protein are
       numbered (from the initiation methionine) on the right.
     ANSWER 24 OF 135 IFIPAT COPYRIGHT 2004 IFI on STN DUPLICATE 13 10094016 IFIPAT; IFIUDB; IFICDB
       POTENTIAL EFFECTOR FOR THE GRB7 FAMILY OF SIGNALLING PROTEINS; NUCLEOTIDE
       SEQUENCES CODING SIGNAL TRANSDUCTION PLYPEPTIDE; FOR USE IN THE DIAGNOSIS
       AND TREATMENT OF CANCERS
       DALY ROGER JOHN (AU); SUTHERLAND ROBERT L (AU)
       Unassigned Or Assigned To Individual (68000)
       US 2002037582
                          A1 20020328
       US 2000-509196
                               20000323
       WO 1998-AU795
                               19980923
       AU 1997-9388
                               19970923
       US 2002037582
                               20020328
       Utility; Patent Application - First Publication
       CHEMICAL
       APPLICATION
       15
        2 Figure(s).
      FIG. 1 provides the nucleotide and amino acid (single letter code)
       sequence of 2.2412. Numbers refer to distances in base pairs.
       Ankyrin-type repeat sequences are underlined. An additional repeat
       sequence is indicated by italics. The stop codon is represented by all asterisk. The original cDNA clone 2. 2412 isolated by the two hybrid screen spans nucleotides 6942664 of this sequence.
     FIG. 2 provides a map of the 2.2412-binding region on ***Grb14*** . A Structure of the deletion constructs used in the analysis. Ga14 DNA-BD fusion constructs encoding full length ***Grb14*** (FL), the Nterminal (N), central region (C) and N-terminal+central region (N+C) were generated in the vector pAS2.1. B. Results of betagalactosidase
       activity assays following transformation of the above plasmids into yeast
       strain Y190 together with the original 2.2412 cDNA clone in pACT-2.
      ANSWER 25 OF 135 CAPLUS COPYRIGHT 2004 ACS ON STN
      2002:10730
                   CAPLUS
      136:49326
      Diagnosis of diseases associated with the immune system using oligomer
      probes to detect cytosine methylation state
      Olek, Alexander; Piepenbrock, Christian; Berlin, Kurt
      Epigenomics A.-G., Germany
      PCT Int. Appl., 32 pp.
      CODEN: PIXXD2
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      German
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                                                                         DATE
     WO 2002000928
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```

EP 2001-953936

20010406

20030115

Α2

Grb14

using two clones L5 and L6, isolated from a human liver cDNA library.

using clone 1 as a probe. The

L6 ΑN

ΤI

IN

PA PΙ

ΑI

FI

DT FS

PRAI

CLMN

GΙ

L6

ΑN

DN

ΤI

IN

PA

S₀

DT

ΙΑ

PΙ

EP 1274865

cDNA sequence was completed

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     ANSWER 26 OF 135
                        USPATFULL on STN
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       2002:315083
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AN
       Nucleic acid sequences associated with baldness
ΤI
       Pritchard, David, Seattle, WA, UNITED STATES
IN
       Burmer, Glenna, Seattle, WA, UNITED STATES
       Brown, Joseph, Seattle, WA, UNITED STATES
       Demas, Vasiliki, Seattle, WA, UNITED STATES
                                 20021128
PΙ
       us 2002177566
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INCL
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L6
     ANSWER 27 OF 135 USPATFULL on STN
ΑN
       2002:181561 USPATFULL
       Process for in vitro selection of high methol producing genotypes
TI
       Khanuja, Suman Preet Singh, Lucknow, INDIA
IN
       Shasany, Ajit Kumar, Lucknow, INDIA
       Dhawan, Sunita, Lucknow, INDÍA
       Darokar, Mahendra Pandurang, Lucknow, INDIA
       Kumar, Tiruppadiripuliyur Ranganathan Santha, Lucknow, INDIA
       Saikia, Dharmendra, Lucknow, INDIA
       Naqui, Arif Ali, Lucknow, INDIA
       Kumar, Sushil, Lucknow, INDIA
       Council of Scientific&Industrial Reaearch, New Delhi, INDIA (non-U.S.
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     ANSWER 28 OF 135
                        WPIDS COPYRIGHT 2004 THOMSON DERWENT on STN
L6
     2002-547451 [58]
ΑN
                         WPIDS
DNC
     C2002-155181
TI
     Treatment or prophylaxis of a subject having a disorder characterized by
     abnormal interaction of Grb7 and a Grb7 ligand, involves administering to
```

a non-phosphorylated peptide to a subject in need of the treatment.

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DC
     B04 D16
IN
     KRAG, D N; OLIGINO, L; PERO, S C
     (UYVE-N) UNIV VERMONT & STATE AGRIC COLLEGE; (KRAG-I) KRAG D N; (OLIG-I)
PA
     OLIGINO L; (PERO-I) PERO S C
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     AU 2002020265 A
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                                                         A61K038-00
     US 2003105000 A1 20030605 (200339)
                                                         A61K038-17
     WO 2002036142 A2 WO 2001-US47400 20011105; AU 2002020265 A AU 2002-20265
ADT
     20011105; US 2003105000 A1 Provisional US 2000-245755P 20001103, US
     2001-13815 20011105
FDT
    AU 2002020265 A Based on WO 2002036142
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                       20001103; US 2001-13815
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     ICM A61K038-00; A61K038-17
IC
L6
     ANSWER 29 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN
     DUPLICATE 14
     2002:492206 BIOSIS
ΑN
     PREV200200492206
DN
     Association of Grb7 with phosphoinositides and its role in the regulation
TT
     of cell migration.
ΑU
     Shen, Tang-Long; Han, Dong Cho; Guan, Jun-Lin [Reprint author]
     Department of Molecular Medicine, Cornell University, Ithaca, NY, 14853,
CS
     USA
     jg19@cornell.edu
     Journal of Biological Chemistry, (August 9, 2002) Vol. 277, No. 32, pp.
SO
     29069-29077. print.
     CODEN: JBCHA3. ISSN: 0021-9258.
     Article
DT
     English
LA
     Entered STN: 18 Sep 2002
ED
     Last Updated on STN: 18 Sep 2002
L6
     ANSWER 30 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN
     DUPLICATE 15
AN
     2002:292126
                   BIOSIS
     PREV200200292126
DN
     Identification of novel non-phosphorylated ligands, which bind selectively
TI
     to the SH2 domain of Grb7.
ΑU
     Pero, Stephanie C.; Oligino, Lyn; Daly, Roger J.; Soden, Amy L.; Liu,
     Chen; Roller, Peter P.; Li, Peng; Krag, David N. [Reprint author]
Department of Surgery, University of Vermont School of Medicine, Given
CS
     Medical Building, Rm. E309, Burlington, VT, 05405, USA
     David.Krag@uvm.edu
SO
     Journal of Biological Chemistry, (April 5, 2002) Vol. 277, No. 14, pp.
     11918-11926. print.
     CODEN: JBCHA3. ISSN: 0021-9258.
DT
     Article
     English
LA
FD
     Entered STN: 15 May 2002
     Last Updated on STN: 15 May 2002
L6
     ANSWER 31 OF 135 CAPLUS COPYRIGHT 2004 ACS on STN
     2002:875636 CAPLUS
ΑN
DN
     138:151170
TI
     Comparative analysis of mutation frequency of coding and non coding short
     mononucleotide repeats in mismatch repair deficient colorectal cancers
     Duval, Alex; Reperant, Maryline; Hamelin, Richard
     INSERM U434, CEPH, Paris, Fr. Oncogene (2002), 21(52), 8062-8066 CODEN: ONCNES; ISSN: 0950-9232
CS
SO
PB
     Nature Publishing Group
     Journal
DT
     English
LA
RE.CNT
        17
               THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
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ΤI
     The adapter protein ZIP binds
                                                      and regulates its inhibitory
```

action on insulin signaling by recruiting protein kinase Czeta.

Cariou, Bertrand; Perdereau, Dominique; Cailliau, Katia; Browaeys-Poly,

ΑU

Edith; Bereziat, Veronique; Vasseur-Cognet, Mireille; Girard, Jean; Burnol, Anne-Francoise [Reprint author] CS Departement d'Endocrinologie, Institut Cochin, CNRS-INSERM-Universite Rene Descartes, 24 Rue du Faubourg Saint-Jacques, 75674, Paris, France burnol@cochin.inserm.fr S₀ Molecular and Cellular Biology, (October, 2002) Vol. 22, No. 20, pp. 6959-6970. print. CODEN: MCEBD4. ISSN: 0270-7306. Article DT English LA Entered STN: 16 Oct 2002 ED Last Updated on STN: 16 Oct 2002 ANSWER 33 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN L6 DUPLICATE 17 ΑN 2002:207358 BIOSIS PREV200200207358 DN Inhibition of insulin receptor catalytic activity by the molecular adapter ***Grb14*** . TI Bereziat, Veronique; Kasus-Jacobi, Anne; Perdereau, Dominique; Cariou, Bertrand; Girard, Jean; Burnol, Anne-Francoise [Reprint author] Endocrinologie et Metabolisme, CNRS UPR 1524, Institut Cochin de Genetique ΑU CS Moleculaire, 24 rue du Faubourg Saint-Jacques, 75674, Paris Cedex, 14, France burnol@cochin.inserm.fr S0 Journal of Biological Chemistry, (February 15, 2002) Vol. 277, No. 7, pp. 4845-4852. print. CODEN: JBCHA3. ISSN: 0021-9258. Article DT English LA ED Entered STN: 20 Mar 2002 Last Updated on STN: 20 Mar 2002 L6 ANSWER 34 OF 135 CAPLUS COPYRIGHT 2004 ACS ON STN ΑN 2003:3572 CAPLUS DN 138:382650 Ontogeny and the possible function of a novel epidermal growth factor-like TI repeat domain-containing protein, NELL2, in the rat brain Kim, Hyun; Ha, Chang Man; Choi, Jungil; Choi, Eun Jung; Jeon, Jongrye; Kim, Changmee; Park, Sang Kyu; Kang, Sang Soo; Kim, Kyungjin; Lee, Byung Department of Anatomy, Brain Korea 21 Biomedical Sciences, Korea University College of Medicine, Seoul, S. Korea Journal of Neurochemistry (2002), 83(6), 1389-1400 CS SO CODEN: JONRA9; ISSN: 0022-3042 PB Blackwell Science Ltd. DT Journal English LA RE.CNT 46 THERE ARE 46 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT L6 ANSWER 35 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN 2002:396091 BIOSIS AN DN PREV200200396091 TI Gene expression profiling of endometrial carcinomas: Identification of molecular biomarkers. Yap, Oi Wah Stephanie [Reprint author]; Zhu, Shirley [Reprint author]; van de Rijn, Matt [Reprint author]; Longacre, Teri [Reprint author]; Teng, Nelson [Reprint author]; Husain, Amreen [Reprint author] CS Stanford University Medical Center, Stanford, CA, USA Proceedings of the American Association for Cancer Research Annual Meeting, (March, 2002) Vol. 43, pp. 746. print. Meeting Info.: 93rd Annual Meeting of the American Association for Cancer Research. San Francisco, California, USA. April 06-10, 2002. ISSN: 0197-016x. Conference; (Meeting)
Conference; Abstract; (Meeting Abstract) DT LA Enalish Entered STN: 24 Jul 2002 ED Last Updated on STN: 24 Jul 2002 L6 ANSWER 36 OF 135 SCISEARCH COPYRIGHT 2004 THOMSON ISI ON STN 2002:207936 SCISEARCH ΑN GΑ The Genuine Article (R) Number: 524UV Comparison of gene expression in old versus young rat hippocampus by cDNA

array

ΑU Cho K S; Choi J G; Ha C M; Son Y J; Choi W S; Lee B J (Reprint) Univ Ulsan, Dept Biol Sci, Ulsan 680749, South Korea (Reprint); Gyeongsang CS Natl Univ, Coll Med, Dept Anat, Chinju 660280, South Korea South Korea CYA NEUROREPORT, (4 MAR 2002) Vol. 13, No. 3, pp. 285-289. Publisher: LIPPINCOTT WILLIAMS & WILKINS, 530 WALNUT ST, PHILADELPHIA, PA S0 19106-3621 USA. ISSN: 0959-4965 Article; Journal DT English IΑ REC Reference Count: 26 *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS* L6 ANSWER 37 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN **DUPLICATE 18** ΑN 2003:124261 BIOSIS PREV200300124261 DN Assignment of backbone 1H, 13C, and 15N resonances of the SH2 domain of TI ***Grb14*** Scharf, Paul J.; Lyons, Barbara A. [Reprint Author] ΑU Department of Biochemistry, College of Medicine, University of Vermont, CS Burlington, VT, 05405, USA blyons@zoo.uvm.edu Journal of Biomolecular NMR, (November 2002) Vol. 24, No. 3, pp. 275-276. SO ISSN: 0925-2738 (ISSN print). Article DT LA English Entered STN: 5 Mar 2003 ED Last Updated on STN: 5 Mar 2003 L6 ANSWER 38 OF 135 SCISEARCH COPYRIGHT 2004 THOMSON ISI ON STN 2002:524711 SCISEARCH ΑN GΑ The Genuine Article (R) Number: 557XP ***Grb14*** TI The atypical PKC-Interacting protein ZIP binds potentiates its inhibitory action on insulin signaling ΑU Cariou B (Reprint); Perdereau D; Cailliau K; Browaeys-Poly E; Bereziat V; Girard J; Burnol A F DIABETES, (JUN 2002) Vol. 51, Supp. [2], pp. A56-A56. MA 228. Publisher: AMER DIABETES ASSOC, 1660 DUKE ST, ALEXANDRIA, VA 22314 USA. S₀ ISSN: 0012-1797. DT Conference; Journal English LA REC Reference Count: 0 L6 ANSWER 39 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN AN 2003:155451 BIOSIS DN PREV200300155451 TI Proteomic Study on the Retinas of RD, RDS and C3B Mice as Well as Reconfirming of the Differentially Expressed Proteins by Using RT-PCT. ΑU Li, D. [Reprint Author]; Zhang, Q. J. Ocular Genetics and Molec Bio, Zhonghsan Ophthalmic Ctr, GuanZhou, China CS ARVO Annual Meeting Abstract Search and Program Planner, (2002) Vol. 2002, pp. Abstract No. 3629. cd-rom. SO Meeting Info.: Annual Meeting of the Association For Research in Vision and Ophthalmology. Fort Lauderdale, Florida, USA. May 05-10, 2002. DT Conference; (Meeting) Conference; Abstract; (Meeting Abstract) ΙΑ English ED Entered STN: 26 Mar 2003 Last Updated on STN: 26 Mar 2003 L6 ANSWER 40 OF 135 CAPLUS COPYRIGHT 2004 ACS on STN 2001:798473 CAPLUS ΑN DN 135:340282 TI Nucleic acid sequences associated with baldness and uses in detecting the likelihood of baldness and for gene therapy IN Pritchard, David; Burmer, Glenna; Brown, Joseph; Demas, Vasiliki Lifespan Biosciences, Inc., USA PΑ S0 PCT Int. Appl., 87 pp. CODEN: PIXXD2 DT Patent LA English FAN.CNT 1 APPLICATION NO. DATE PATENT NO. KIND DATE

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wo 2001-US12184 20010413
PΙ
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L6
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     2001:390236
                   BIOSIS
AN
     PREV200100390236
DN
     Identification of a novel human tankyrase through its interaction with the
TI
                         ***Grb14***
     adaptor protein
     Lyons, Ruth J.; Deane, Roisin; Lynch, Danielle K.; Ye, Zheng-Sheng
     Jeffrey; Sanderson, Georgina M.; Eyre, Helen J.; Sutherland, Grant R.;
     Daly, Roger J. [Reprint author]
     Cancer Research Program, Garvan Institute of Medical Research, St.
CS
     Vincent's Hospital, Sydney, NSW, 2010, Australia
     r.daly@garvan.org.au
     Journal of Biological Chemistry, (May 18, 2001) Vol. 276, No. 20, pp.
S0
     17172-17180. print.
     CODEN: JBCHA3. ISSN: 0021-9258.
     Article
DT
     English
LA
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ED
      Entered STN: 15 Aug 2001
      Last Updated on STN: 23 Feb 2002
     ANSWER 42 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN
L6
     DUPLICATE 20
      2001:514026
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AN
DN
      PREV200100514026
     The Grb7 family proteins: Structure, interactions with other signaling
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     molecules and potential cellular functions.
      Han, Dong Cho; Shen, Tang-Long; Guan, Jun-Lin [Reprint author]
ΔH
      Cancer Biology Laboratories, Department of Molecular Medicine, Cornell
CS
      University, Ithaca, NY, 14853, USA
jg19@cornell.edu
      Oncogene, (1 October, 2001) Vol. 20, No. 44, pp. 6315-6321. print.
S0
      CODEN: ONCNES. ISSN: 0950-9232.
DT
      Article
      General Review; (Literature Review)
      English
LA
      Entered STN: 7 Nov 2001
ED
      Last Updated on STN: 23 Feb 2002
      ANSWER 43 OF 135 CAPLUS COPYRIGHT 2004 ACS on STN
L6
ΑN
      2001:634713
                   CAPLUS
      135:342315
DN
      Extensive characterization of genetic alterations in a series of human
TI
      colorectal cancer cell lines
      Gayet, Jacqueline; Zhou, Xiao-Ping; Duval, Alex; Rolland, Sandra; Hoang,
ΑU
      Jean-Marc; Cottu, Paul; Hamelin, Richard
      INSERM U434 - CEPH, Paris, 75010, Fr.
CS
                                  5025-5032
      Oncogene (2001), 20(36),
S<sub>0</sub>
      CODEN: ONCNES; ISSN: 0950-9232
      Nature Publishing Group
PR
      Journal
DT
      English
LA
                THERE ARE 63 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 63
                ALL CITATIONS AVAILABLE IN THE RE FORMAT
      ANSWER 44 OF 135 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 21
L6
      2001:170169
AN
      135:270832
DN
      Evolution of instability at coding and non-coding repeat sequences in
TI
```

human MSI-H colorectal cancers

```
Duval, Alex; Rolland, Sandra; Compoint, Aurore; Tubacher, Emmanuel;
ΑU
     Iacopetta, Barry; Thomas, Gilles; Hamelin, Richard
     INSERM U434, CEPH, Paris, 75010, Fr.
Human Molecular Genetics (2001), 10(5), 513-518
CS
50
      CODEN: HMGEE5; ISSN: 0964-6906
PB
      Oxford University Press
      Journal
DT
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                THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS RECORD
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L6
      DUPLICATE 22
      2001:209006
                    BIOSIS
ΑN
      PREV200100209006
DN
      The BPS domain of Grb10 inhibits the catalytic activity of the insulin and
TT
      IGF1 receptors.
     Stein, Evan G.; Gustafson, Thomas A.; Hubbard, Stevan R. [Reprint author] Department of Pharmacology, Skirball Institute of Biomolecular Medicine, New York University School of Medicine, 540 First Avenue, New York, NY,
CS
      10016, USA
      hubbard@tallis.med.nyu.edu
      FEBS Letters, (30 March, 2001) Vol. 493, No. 2-3, pp. 106-111. print.
$0
      CODEN: FEBLAL. ISSN: 0014-5793.
      Article
DT
      English
LA
ED
      Entered STN: 2 May 2001
      Last Updated on STN: 18 Feb 2002
      ANSWER 46 OF 135 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN
L6
      2002:69396 BIOSIS
ΑN
      PREV200200069396
DN
      Inhibition of insulin receptors tyrosine kinase activity by the molecular
TI
                  ***Grb14***
      Bereziat, V. [Reprint author]; Kasus-Jacobi, A. [Reprint author];
      Perdereau, D. [Reprint author]; Girard, J. [Reprint author]; Burnol, A.-F.
      [Reprint author]
     CNRS UPR1524, ICGM, 9 rue Jules Hetzel, 92190, Meudon, France Biochemical Society Transactions, (2001) Vol. 29, No. 3, pp. A69. print. Meeting Info.: 673rd Bristol Meeting of the Biochemical Society. Bristol, London, UK. April 10-12, 2000. Biochemical Society.
CS
SO
      CODEN: BCSTB5. ISSN: 0300-5127.
      Conference; (Meeting)
Conference; Abstract; (Meeting Abstract)
DT
      English
LA
      Entered STN: 16 Jan 2002
ED
      Last Updated on STN: 25 Feb 2002
      ANSWER 47 OF 135 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 23
L6
      2001:131521 CAPLUS
ΑN
DN
      134:202914
      Human growth factor receptor bound 14 binds the activated insulin receptor
TI
      and alters the insulin-stimulated tyrosine phosphorylation levels of
      multiple proteins
      Hemming, Richard; Agatep, Ronald; Badiani, Ketan; Wyant, Kerrie; Arthur.
ΑU
      Gilbert; Gietz, R. Daniel; Triggs-Raine, Barbara
      Department of Biochemistry & Medical Genetics, University of Manitoba,
CS
      Winnipeg, MB, R3E 0W3, Can.
      Biochemistry and Cell Biology (2001), 79(1), 21-32
SO
      CODEN: BCBIEQ; ISSN: 0829-8211
PB
      National Research Council of Canada
DT
      Journal
      English
LA
RE.CNT
        46
                THERE ARE 46 CITED REFERENCES AVAILABLE FOR THIS RECORD
                ALL CITATIONS AVAILABLE IN THE RE FORMAT
L6
      ANSWER 48 OF 135 DISSABS COPYRIGHT (C) 2004 ProQuest Information and
      Learning Company; All Rights Reserved on STN
ΑN
      2001:8470
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      Jones, Nina [Ph.D.]; Dumont, Daniel J. [adviser]
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      University of Toronto (Canada) (0779)
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                              Order Number: AAIMQ51679
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                              interacting protein
      Agatep, Ronald [M.Sc.]; Gietz, R. D. [adviser]
The University of Manitoba (Canada) (0303)
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                       proteins for screening compounds capable of modulating
      insulin receptor tyrosine kinase activity
      Burnol, Anne-Francoise; Perdereau, Dominique; Kasus-Jacobi, Anne;
 IN
      Bereziat, Veronique; Girard, Jean
 PΑ
      Centre National De La Recherche Scientifique-CNRS, Fr.
      PCT Int. Appl., 46 pp.
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TI
                 ***Grb14*** . Characterization of a new receptor binding
      protein
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      Reilly, John F.; Mickey, Gregory; Maher, Pamela A. [Reprint author]
     Dept. of Cell Biology, The Scripps Research Institute, 10550 N. Torrey
CS
     Pines Rd., CAL-3, La Jolla, CA, 92037, USA
Journal of Biological Chemistry, (March 17, 2000) Vol. 275, No. 11, pp.
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      CODEN: JBCHA3. ISSN: 0021-9258.
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     for two of its binding domains, PIR and SH2.
     Kasus-Jacobi, Anne; Bereziat, Veronique; Perdereau, Dominique; Girard, Jean; Burnol, Anne-Francoise [Reprint author] Endocrinologie Metabolisme et Developpement, CNRS, UPR 1524, 9 Rue Jules
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      Morrione, Andrea [Reprint author]
ΑU
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CS
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SO
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      ISSN: 1107-3756.
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      Last Updated on STN: 7 Jan 2002
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      Potential effector protein for the Grb7 family of signaling proteins
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      Daly, Roger John; Sutherland, Robert Lyndsay
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      CODEN: PIXXD2
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ΑU
     author]
CS
     Sunnybrook and Women's College Health Sciences Centre, 2075 Bayview Ave..
     Research Bldg., S-227, Toronto, ON, M4N 3M5, Canada
SO
      Journal of Biological Chemistry, (oct. 22, 1999) Vol. 274, No. 43, pp.
      30896-30905. print.
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Dep. Cell Biol., Scripps Res. Inst., La Jolla, CA 92037, USA
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                        Univ. Texas Health Sci. Cent., 7703 Floyd Curl Dr., San
      Antonio, TX 78284-7764, USA
      Journal of Biological Chemistry, (Nov. 14, 1997) vol. 272, No. 46, pp.
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     the Src homology 2 domain of Grb7.
     Janes, Peter W.; Lackmann, Martin; Church, W. Bret; Sanderson, Georgina M.; Sutherland, Robert L.; Daly, Roger J. [Reprint author]
Cancer Res. Program, Garvan Inst. Med. Res., St. Vincent's Hosp., Sydney,
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CS
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      Journal of Biological Chemistry, (1997) Vol. 272, No. 13, pp. 8490-8497.
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TI
     Human GRB-IR-beta/GRB10: Splice variants of an insulin and growth factor
     receptor-binding protein with PH and SH2 domains.
     Frantz, J. Daniel; Giorgetti-Peraldi, Sophie; Ottinger, Elizabeth A.;
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     Shoelson, Steven E. [Reprint author]
     Joslin Diabetes Cent., One Joslin Place, Boston, MA 02215, USA Journal of Biological Chemistry, (1997) Vol. 272, No. 5, pp. 2659-2667.
SO
     CODEN: JBCHA3. ISSN: 0021-9258.
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     GDU: a new target for the erbB family of protein tyrosine kinases and a
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     Daly, Roger John; Sutherland, Robert Lyndsay
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AN
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      Cloning and characterization of ***GRB14***
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      GRB7 gene family.
ΑU
      Daly, Roger J. [Reprint author]; Sanderson, Georgina M.; Janes, Peter W.;
      Sutherland, Robert L.
Cancer Biol. Div., Garvan Inst. Med. Res., St. Vincent's Hosp., Sydney,
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      Assignment of the human
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      Baker, Elizabeth; Sutherland, Grant R.; Sutherland, Robert L.; Daly, Roger
ΑU
      J. [Reprint author]
      Cancer Biol. Div., Garvan Inst. Med. Res., St. Vincent's Hosp., Sydney,
CS
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AN

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DGENE

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A new signalling protein designated GDU related to erbB receptor targets
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GenBank ACC. NO. (GBN): AY419444
GenBank VERSION (VER): AY419444
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COMMENT:
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                          Clark,A.G.; Glanowski,S.; Nielson,R.; Thomas,P.;
Kejariwal,A.; Todd,M.A.; Tanenbaum,D.M.; Civello,D.R.;
   AUTHOR (AU):
                          Lu, F.; Murphy, B.; Ferriera, S.; Wang, G.; Zheng, X.H.;
                          White, T.J.; Sninsky, J.J.; Adams, M.D.; Cargill, M. Inferring nonneutral evolution from human-chimp-mouse
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Science, 302 (5652), 1960-1963 (2003)
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                          White, T.J.; Sninsky, J.J.; Adams, M.D.; Cargill, M.
   TITLE (TI):
                          Direct Submission
                          Submitted (16-NOV-2003) Celera Genomics, 45 West Gude
   JOURNAL (SO):
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L6
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LOCUS (LOC):
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REFERENCE:
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                        Clark, A.G.; Glanowski, S.; Nielson, R.; Thomas, P.; Kejariwal, A.; Todd, M.A.; Tanenbaum, D.M.; Civello, D.R.; Lu, F.; Murphy, B.; Ferriera, S.; Wang, G.; Zheng, X.H.; White, T.J.; Sninsky, J.J.; Adams, M.D.; Cargill, M. Inferring nonneutral evolution from human-chimp-mouse
   AUTHOR (AU):
   TITLE (TI):
                        orthologous gene trios
Science, 302 (5652), 1960-1963 (2003)
   JOURNAL (SO):
REFERENCE:
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L6
      ANSWER 75 OF 135
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DATE (DATE):
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L6
       ANSWER 76 OF 135
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DATE (DATE):
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COMMENT:
       Contact: MGC help desk
       Email: cgapbs-r@mail.nih.gov
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       CDNA Library Preparation: Life Technologies, Inc.
       cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
       DNA Sequencing by: National Institutes of Health Intramural
       Sequencing Center (NISC),
       Gaithersburg, Maryland;
       Web site: http://www.nisc.nih.gov/
       Contact: nisc_mgc@nhgri.nih.gov
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    TITLE (TI):
                                  Generation and initial analysis of more than 15,000
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    JOURNAL (SO):
                                  Proc. Natl. Acad. Sci. U.S.A., 99 (26), 16899-16903
                                  (2002)
    OTHER SOURCE (OS):
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REFERENCE:
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    TITLE (TI):
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JOURNAL (SO):

Submitted (09-JUN-2003) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA

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    TITLE (TI):
                                 Direct Submission
                                 Submitted (25-MAR-2002) Takao Isogai, Helix Research
    JOURNAL (SO):
                                 Institute, Genomics Laboratory; 1532-3 Yana, Kisarazu,
                                 Chiba 292-0812, Japan (E-mail:genomics@hri.co.jp,
                                 Tel:81-438-52-3975, Fax:81-438-52-3986)
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L6
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LOCUS (LOC):
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MOLECULE TYPE (CI):
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DIVISION CODE (CI):
DATE (DATE):
                              16 Apr 2003
DEFINITION (DEF):
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                              Mus musculus (house mouse)
SOURCE:
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NUCLEIC ACID COUNT (NA): 277 a
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COMMENT:
      Contact: MGC help desk
      Email: cgapbs-r@mail.nih.gov
      Tissue Procurement: Gilbert Smith, Ph.D.
      CDNA Library Preparation: Life Technologies, Inc.
      CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
      DNA Sequencing by: Baylor College of Medicine Human Genome
      Sequencing Center
      Center code: BCM-HGSC
      web site: http://www.hgsc.bcm.tmc.edu/cdna/
      Contact: amg@bcm.tmc.edu
      Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Loulseged, H.,
      Kowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Nanavati,
      A.N., Gibbs, R.A.
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   AUTHOR (AU):
                              Strausberg, R.L.; Feingold, E.A.; Grouse, L.H.;
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Dickson,M.C.; Rodriguez,A.C.; Grimwood,J.; Schmutz,J.;

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                           full-length human and mouse cDNA sequences
   JOURNAL (SO):
                           Proc. Natl. Acad. Sci. U.S.A., 99 (26), 16899-16903
                           (2002)
   OTHER SOURCE (OS):
                           CA 138:131969
REFERENCE:
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   AUTHOR (AU);
                           Strausberg, R.
   TITLE (TI):
                           Direct Submission
                           Submitted (18-JAN-2002) National Institutes of Health,
    JOURNAL (SO):
                           Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive,
                           Room 11A03, Bethesda, MD 20892-2590, USA
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DIVISION CODE (CI):

Primates

Myers, R.M.; Butterfield, Y.S.; Krzywinski, M.I.;

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14 May 2001
DATE (DATE):
DEFINITION (DEF):
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   AUTHOR (AU):
                          Lyons, R.J.; Deane, R.; Lynch, D.K.; Ye, Z.S.;
                          Sanderson, G.M.; Eyre, H.J.; Sutherland, G.R.; Daly, R.J.
                          Identification of a novel human tankyrase through its interaction with the adaptor protein ***Grb14***

J. Biol. Chem., 276 (20), 17172-17180 (2001)
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                          Lyons, R.J.; Deane, R.; Lynch, D.K.; Ye, Z.-S.J.;
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                          Sanderson,G.M.; Eyre,H.J.; Sutherland,G.R.; Daly,R.J.
   TITLE (TI):
                          Direct Submission
                          Submitted (17-DEC-2000) Cancer Research, Garvan Institute of Medical Research, 384 Victoria St.,
   JOURNAL (SO):
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L6 ANSWER 80 OF 135 GENBANK.RTM. COPYRIGHT 2004 on STN

LOCUS (LOC):

GenBank ACC. NO. (GBN): BB560608

GenBank VERSION (VER): BB560608.1

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SEQUENCE LENGTH (SQL): 323

MOLECULE TYPE (CI): mRNA; linear

MOLECULE TYPE (CI): mRNA; linear Expressed sequence tag DATE (DATE): 1 Aug 2000

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COMMENT:
       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic
       Sciences Center(GSC), Yokohama Institute
       The Institute of Physical and Chemical Research (RIKEN)
       1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
       Tel: 81-45-503-9222
       Fax: 81-45-503-9216
       Email: genome-res@gsc.riken.go.jp,
       URL:http://genome.gsc.riken.go.jp/
       Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Thermostabilization and thermoactivation of thermolabile enzymes by
       trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.,
       Tomarú,Yí, Carninci,Pí, Shibatá,Yí, Ozawa,Ý., Muramatsu,M., Okázaki
       ,Y. and Hayashizaki,Y.
        Automated filtration-based high-throughput plasmid preparation
       system. Genome Res. 9 (5), 463-470 (1999)
        Carninci, P. and Hayashizaki, Y.
        High-efficiency full-length cDNA cloning, Methods Enzymol, 303,
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        Please visit our web site (http://genome.rtc.riken.go.jp) for
       further details.
REFERENCE:
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                                 Konno,H.; Aizawa,K.; Akahira,S.; Akiyama,J.;
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Owa,C.; Saito,H.; Sakai,C.; Sato,K.; Shihata,K.;
    AUTHOR (AU):
                                  Owa, C.; Saito, H.; Sakai, C.; Sato, K.; Shibata, K.
                                  Shibata, Y.,; Shigemoto, Y.; Shinagawa, A.; Shiraki, T.;
                                 Sogabe, Y.; Sugahara, Y., Suzuki, H.; Suzuki, H.; Tagawa, A.; Takahashi, F.; Tominaga, N.; Toya, T.a; Tsunoda, Y.; Watahiki, A.; Watanabe, S.; Yamamura, T.;
                                  Yamanaká,I., Yano,R.H; Yasunishi,Á.; Yokota,T.;
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    TITLE (TI):
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    JOURNAL (SO):
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                                                         Encyclopedia Project of Genome
                                                         Exploration Research Group in
                                                         Riken Genomic Sciences Center and
                                                         Genome Science Laboratory in
                                                         RIKEN. Division of Experimental
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Animal Research in Riken

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L6
      ANSWER 81 OF 135
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LOCUS (LOC):
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GenBank VERSION (VER):
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SEQUENCE LENGTH (SQL):
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DIVISION CODE (CI):
                              Expressed sequence tag
DATE (DATE):
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DEFINITION (DEF):
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NUCLEIC ACID COUNT (NA): 100 a
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COMMENT:
      Contact: Yoshihide Hayashizaki
      Laboratory for Genome Exploration Research Group, RIKEN Genomic
      Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
      Fax: 81-45-503-9216
      Email: genome-res@gsc.riken.go.jp
      URL:http://genome.gsc.riken.go.jp/
      Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki
      N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
      Thermostabilization and thermoactivation of thermolabile enzymes by
      trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
      Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.
       Automated filtration-based high-throughput plasmid preparation
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       Carninci, P. and Hayashizaki, Y.
       High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
      19-44 (1999)
       Please visit our web site (http://genome.rtc.riken.go.jp) for
      further details.
REFERENCE:
                                 (bases 1 to 313)
   AUTHOR (AU):
                              Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.; Arakawa, T.; Carninci, P.; Endo, T.; Fukuda, S.;
                              Fukunishi,Y.; Hara,A.; Hayatsu,N.; Hirozane,Ť.;
Hori,F.; Ishii,Y.; Ishikawa,J.; Ishikawa,T.; Itoh,M.;
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Izawa, M.; Kadota, K.; Kagawa, I.; Kai, C.; Kawai, J.;
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                                                 RIKEN. Division of Experimental
                                                Animal Research in Riken
                                                 contributed to prepare mouse
                                                 tissues. 1st strand cDNA was
                                                 primed with a primer [5
                                                 GAGAGAGAGCGGCCGCAACTCGAGTTTTTTTT
                                                TTTTTTTVN 3'], cDNA was prepared by using trehalose
                                                thermo-activated reverse
transcriptase and subsequently
enriched for full-length by
                                                 cap-trapper. Second strand cDNA
                                                was prepared with the primer
                                                 adapter of sequence [5]
                                                 GAGAGAGATTCTCGAGTTAATTAAATTAATCC
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     ANSWER 82 OF 135
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adapter rGrb14 (***Grb14***) mRNA, mRNA sequence.

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FEATURES (FEAT):

Feature Key

SEQUENCE (SEQ):

LOCUS (LOC):

DATE (DATE):

MOLECULE TYPE (CI):

DIVISION CODE (CI):

DEFINITION (DEF):

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source

JOURNAL (SO):

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NUCLEIC ACID COUNT (NA): 117 a
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COMMENT:
        Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic
       Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
       Fax: 81-45-503-9216
       Email: genome-res@gsc.riken.go.jp,
       URL:http://genome.gsc.riken.go.jp/
       Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki
        ,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
       Thermostabilization and thermoactivation of thermolabile enzymes by
       trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520–524 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.
         Automated filtration-based high-throughput plasmid preparation
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High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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         Please visit our web site (http://genome.rtc.riken.go.jp) for
       further details.
REFERENCE:
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Konno,H.; Aizawa,K.; Akahira,S.; Akiyama,J.;
Arakawa,T.; Carninci,P.; Endo,T.; Fukuda,S.;
Fukunishi,Y.; Hara,A.; Hayatsu,N.; Hirozane,T.;
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    AUTHOR (AU):
                                     Sogabe, Y.; Sugahara, Y., Suzuki, H.; Suzuki, H.; Tagawa, A.; Takahashi, F.; Tominaga, N.; Toya, T.a;
                                     Tsunoda, Y.; Watahiki, A.; Watanabe, S.; Yamamura, T.;
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                                                              contributed to prepare mouse
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primed with a primer [5

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     ANSWER 83 OF 135
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L6
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COMMENT:
     Contact: Yoshihide Hayashizaki
     Laboratory for Genome Exploration Research Group, RIKEN Genomic
     Sciences Center(GSC), Yokohama Institute
     The Institute of Physical and Chemical Research (RIKEN)
     1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
     Fax: 81-45-503-9216
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     Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki
      ,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
     Thermostabilization and thermoactivation of thermolabile enzymes by
     trehalose and its application for the synthesis of full length
     cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.
     Tomaru,Y., Carninci,P., Shibata,Y., Ozawa,Y., Muramatsu,M., Okazaki
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      Automated filtration-based high-throughput plasmid preparation
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      High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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REFERENCE:
                             (bases 1 to 319)
                           Konno,H.; Aizawa,K.; Akahira,S.; Akiyama,J.; Arakawa,T.; Carninci,P.; Endo,T.; Fukuda,S.;
   AUTHOR (AU):
                           Fukunishi,Y.; Hara,A.; Hayatsu,N.; Hirozane,T.;
Hori,F.; Ishii,Y.; Ishikawa,J.; Ishikawa,T.; Itoh,M.;
Izawa,M.; Kadota,K.; Kagawa,I.; Kai,C.; Kawai,J.;
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Mizuno,Y.; Nakamura,M.; Oda,H.; Okazaki,Y., Ono,T.y; Owa,C.; Saito,H.; Sakai,C.; Sato,K.; Shibata,K.; Shibata,Y.,; Shigemoto,Y.; Shinagawa,A.; Shiraki,T.;
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                                                Genome Science Laboratory in
                                                RIKEN. Division of Experimental
                                                Animal Research in Riken
                                                contributed to prepare mouse
                                                tissues. 1st strand cDNA was
                                                primed with a primer [5
                                                GAGAGAGAGCGGCCGCAACTCGAGTTTTTTTT
                                                TTTTTTTVN 3'], cDNA was prepared
                                                by using trehalose
                                                thermo-activated reverse
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                                                enriched for full-length by
                                                cap-trapper. Second strand cDNA
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LOCUS (LOC):
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L6

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       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic
       Sciences Center(GSC), Yokohama Institute
       The Institute of Physical and Chemical Research (RIKEN)
       1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
       Tel: 81-45-503-9222
       Fax: 81-45-503-9216
       Email: genome-res@gsc.riken.go.jp,
       URL:http://genome.gsc.riken.go.jp/
      Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Thermostabilization and thermoactivation of thermolabile enzymes by
      trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.
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        High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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        Please visit our web site (http://genome.rtc.riken.go.jp) for
       further details.
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Fukunishi,Y.; Hara,A.; Hayatsu,N.; Hirozane,T.;
    AUTHOR (AU):
                                Hori, F.; Ishii, Y.; Ishikawa, J.; Ishikawa, T.; Itoh, M.;
                                Izawa,M.; Kadota,K.; Kagawa,I.; Kai,C.; Kawai,J.;
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Encyclopedia Project of Genome
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                                                      RIKEN. Division of Experimental
                                                      Animal Research in Riken
                                                      contributed to prepare mouse
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SOURCE:
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NUCLEIC ACID COUNT (NA): 106 a
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COMMENT:
       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic
       Sciences Center(GSC), Yokohama Institute
      The Institute of Physical and Chemical Research (RIKEN)
      1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222 Fax: 81-45-503-9216
       Email: genome-res@gsc.riken.go.jp,
      URL:http://genome.gsc.riken.go.jp/
       Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki
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       Thermostabilization and thermoactivation of thermolabile enzymes by
      trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.
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        Carninci, P. and Hayashizaki, Y.
        High-efficiency full-length cDNA cloning. Methods Enzymol. 303.
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        Please Visit our web site (http://genome.rtc.riken.go.jp) for
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REFERENCE:
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                                 Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.; Arakawa, T.; Carninci, P.; Endo, T.; Fukuda, S.; Fukunishi, Y.; Hara, A.; Hayatsu, N.; Hirozane, T.;
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Animal Research in Riken
                                                     contributed to prepare mouse
                                                     tissues. 1st strand cDNA was
                                                     primed with a primer [5
                                                     GAGAGAGAGAGCGCCCGCAACTCGAGTTTTTTTT
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       ANSWER 86 OF 135
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DATE (DATE):
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NUCLEIC ACID COUNT (NA): 87 a 45 c 38 g
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Contact: Yoshihide Hayashizaki
      Laboratory for Genome Exploration Research Group, RIKEN Genomic
      Sciences Center(GSC), Yokohama Institute
      The Institute of Physical and Chemical Research (RIKEN)
      1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
      Tel: 81-45-503-9222
      Fax: 81-45-503-9216
      Email: genome-res@gsc.riken.go.jp
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      Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y. Thermostabilization and thermoactivation of thermolabile enzymes by
      trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.
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        Automated filtration-based high-throughput plasmid preparation
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High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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                                                          GAGAGAGAGAGGATCCAAGAGCTCTTTTTTTT
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L6
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COMMENT:
      Contact: Yoshihide Hayashizaki
      Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
      1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
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      Email: genome-res@gsc.riken.go.jp,
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      Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki
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      Thermostabilization and thermoactivation of thermolabile enzymes by
      trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki
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       Please visit our web site (http://genome.rtc.riken.go.jp) for
      further details.
REFERENCE:
                                 (bases 1 to 284)
                             Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.; Arakawa, T.; Carninci, P.; Endo, T.; Fukuda, S.;
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                             Fukunishi, Y.; Hara, A.; Hayatsu, N.; Hirozane, T.;
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Animal Research in Riken
                                                 contributed to prepare mouse
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enriched for full-length by
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round of normalization to Rot =
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           Contact: Yoshihide Hayashizaki
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           The Institute of Physical and Chemical Research (RIKEN)
           1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
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Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
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             High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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tissues. 1st strand cDNA was

house mouse.

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                                         Lambda FLC I. Cloning sites, 5' end: SalI; 3' end: BamHI"
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                                        61 g
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  Laboratory for Genome Exploration Research Group, RIKEN Genomic
  Sciences Center(GSC), Yokohama Institute
  The Institute of Physical and Chemical Research (RIKEN)
  1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
  Email: genome-res@gsc.riken.go.jp,
  URL:http://genome.gsc.riken.go.jp/
  Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki
  N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
  Thermostabilization and thermoactivation of thermolabile enzymes by
  trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
  Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki
   Automated filtration-based high-throughput plasmid preparation
  system. Genome Res. 9 (5), 463-470 (1999)
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   Please visit our web site (http://genome.rtc.riken.go.jp) for
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DATE (DATE):

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Fax: 81-45-503-9216

,Y. and Hayashizaki,Y.

19-44 (1999)

AUTHOR (AU):

REFERENCE:

further details.

Carninci, P. and Hayashizaki, Y.

Contact: Yoshihide Hayashizaki

BB319216

311

278502-24-2

mRNA; linear

11 Jul 2000

house mouse.

Mus musculus

69 c

Konno,H.; Aizawa,K.; Akahira,S.; Akiyama,J.;
Arakawa,T.; Carninci,P.; Endo,T.; Fukuda,S.;

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                                                         RIKEN. Division of Experimental Animal Research in Riken
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                                                         GAGAGAGAAGGATCCAAGAGCTCTTTTTTTT
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ANSWER 90 OF 135

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LOCUS (LOC):

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TITLE (TI):

FEATURES (FEAT): Feature Key

source

JOURNAL (SO):

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COMMENT:
       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic
       Sciences Center(GSC), Yokohama Institute
       The Institute of Physical and Chemical Research (RIKEN)
      1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222 Fax: 81-45-503-9216
       Email: genome-res@gsc.riken.go.jp,
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       Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
       Thermostabilization and thermoactivation of thermolabile enzymes by
       trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
        Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.
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        Automated filtration-based high-throughput plasmid preparation
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        Carninci,P. and Hayashizaki,Y.
        High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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        Please visit our web site (http://genome.rtc.riken.go.jp) for
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REFERENCE:
                                Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.;
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    AUTHOR (AU):
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```

Genome Science Laboratory in

RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5 GAGAGAGAAGGATCCAAGAGCTCTTTTTTTT TTTTTTTVN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to Rot = 20.0 and subtraction to Rot = 459.0. Second strand cDNA was prepared with the primer adapter of sequence [5'GAGAGAGAGATTCTCGAGTTAATTAAATTAA TCCCCCCCCCCC 3']. cDNA was cleaved with XhoI and BamHI. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FLC I. ~Retina RNA was provided by Stefano Gustincich, Department of Neurobiology Harvard Medical School, 220 Longwood Ave., Boston, MA02115, USA, whose assistance we gratefully acknowledge

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L6
     ANSWER 91 OF 135
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COMMENT:
     Contact: Yoshihide Hayashizaki
     Laboratory for Genome Exploration Research Group, RIKEN Genomic
     Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
     1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
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     Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki
      N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
     Thermostabilization and thermoactivation of thermolabile enzymes by
     trehalose and its application for the synthesis of full length
     cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
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Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.
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High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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       Please visit our web site (http://genome.rtc.riken.go.jp) for
      further details.
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                               Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.;
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                                                    provided by Stefano Gustincich,
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Department of Neurobiology, Harvard Medical School, 220 Longwood Ave., Boston, MA02115,

USA, whose assistance we

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       Laboratory for Genome Exploration Research Group, RIKEN Genomic
       Sciences Center(GSC), Yokohama Institute
       The Institute of Physical and Chemical Research (RIKEN)
       1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
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       Fax: 81-45-503-9216
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       Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Thermostabilization and thermoactivation of thermolabile enzymes by
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Qualifier

Location

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Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
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Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsc.riken.go.jp,

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URL:http://genome.gsc.riken.go.jp/
      Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki
       ,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
      Thermostabilization and thermoactivation of thermolabile enzymes by
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        Carninci, P. and Hayashizaki, Y.
        High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
      19-44 (1999)
        Please visit our web site (http://genome.rtc.riken.go.jp) for
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                                1 (bases 1 to 289)
REFERENCE:
                                Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.; Arakawa, T.; Carninci, P.; Endo, T.; Fukuda, S.; Fukunishi, Y.; Hara, A.; Hayatsu, N.; Hirozane, T.; Hori, F.; Ishii, Y.; Ishikawa, J.; Ishikawa, T.; Itoh, M.;
   AUTHOR (AU):
                                Izawa, M.; Kadota, K.; Kagawa, I.; Kai, C.; Kawai, J.;
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                                                      GAGAGAGAAGGATCCAAGAGCTCTTTTTTTT
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                                                      thermo-activated reverse
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modified pBluescript KS(+) after

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NUCLEIC ACID COUNT (NA): 87 a
COMMENT:
       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic
       Sciences Center(GSC), Yokohama Institute
       The Institute of Physical and Chemical Research (RIKEN)
       1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
       Tel: 81-45-503-9222
       Fax: 81-45-503-9216
       Email: genome-res@gsc.riken.go.jp,
       URL:http://genome.gsc.riken.go.jp/
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       Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki
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        Carninci, P. and Hayashizaki, Y.
        High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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Email: genome-res@gsc.riken.go.jp,
URL:http://genome.gsc.riken.go.jp/

Fax: 81-45-503-9216

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Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki
       ,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
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REFERENCE:
                                  Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.;
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                                                           primed with a primer [5
                                                           GAGAGAGAGAGGATCCAAGAGCTCTTTTTTTT
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                                                           by using trehalose
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                                                           round of normalization to Rot =
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modified pBluescript KS(+) after bulk excision from Lambda FLC I. "

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       ANSWER 96 OF 135
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NUCLEIC ACID COUNT (NA): 79 a
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COMMENT:
        Contact: Yoshihide Hayashizaki
        Laboratory for Genome Exploration Research Group, RIKEN Genomic
        Sciences Center(GSC), Yokohama Institute
        The Institute of Physical and Chemical Research (RIKEN)
        1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
        Tel: 81-45-503-9222
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        URL:http://genome.gsc.riken.go.jp/
       Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
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                                               COPYRIGHT 2004 on STN
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Sciurognathi; Muridae; Murinae; Mus
NUCLEIC ACID COUNT (NA): 108 a
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COMMENT:
      Contact: Yoshihide Hayashizaki
      Laboratory for Genome Exploration Research Group, RIKEN Genomic
      Sciences Center(GSC), Yokohama Institute
      The Institute of Physical and Chemical Research (RIKEN)
      1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
      Tel: 81-45-503-9222
      Fax: 81-45-503-9216
      Email: genome-res@gsc.riken.go.jp,
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URL:http://genome.gsc.riken.go.jp/

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Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki
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       Thermostabilization and thermoactivation of thermolabile enzymes by
       trehalose and its application for the synthesis of full length CDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki
       ,Y. and Hayashizaki,Y.
        Automated filtration-based high-throughput plasmid preparation
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REFERENCE:
                                      (bases 1 to 312)
                                  Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.; Arakawa, T.; Carninci, P.; Endo, T.; Fukuda, S.; Fukunishi, Y.; Hara, A.; Hayatsu, N.; Hirozane, T.; Hori, F.; Ishii, Y.; Ishikawa, J.; Ishikawa, T.; Itoh, M.; Izawa, M.; Kadota, K.; Kagawa, I.; Kai, C.; Kawai, J.; Kikuchi, N.; Kiyosawa, H.; Kojima, Y.; Kondo, S.; Koya, S.; Kurihara, C.; Kusakabe, M., Matsuyama, T.; Miki, R.; Mizuno, Y.; Nakamura, M.; Oda, H.; Okazaki, Y., Ono, T.y; Owa, C.; Saito, H.; Sakai, C.; Saito, K.; Shihata, K.;
    AUTHOR (AU):
                                  Owa, C.; Saito, H.; Sakai, C.; Sato, K.; Shibata, K.; Shibata, Y.,; Shigemoto, Y.; Shinagawa, A.; Shiraki, T.;
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Animal Research in Riken
                                                          contributed to prepare mouse
                                                          tissues. 1st strand cDNA was
                                                          primed with a primer [5
                                                          GAGAGAGAGAAGGATCCAAGAGCTCTTTTTTTT
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                                                          by using trehalose
                                                          thermo-activated reverse
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round of normalization to Rot =
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                                                          prepared with the primer adapter
                                                          of sequence [5'
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modified pBluescript KS(+) after

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           ANSWER 98 OF 135
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L6
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NUCLEIC ACID COUNT (NA): 89 a
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COMMENT:
            Contact: Yoshihide Hayashizaki
           Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
            1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
           Tel: 81-45-503-9222
            Fax: 81-45-503-9216
            Email: genome-res@gsc.riken.go.jp,
           URL:http://genome.gsc.riken.go.jp/
           Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Thermostabilization and thermoactivation of thermolabile enzymes by
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            ,Y. and Hayashizaki,Y.
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             Carninci, P. and Hayashizaki, Y.
             High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
            19-44 (1999)
              Please visit our web site (http://genome.rtc.riken.go.jp) for
            further details.
REFERENCE:
                                                                (bases 1 to 235)
                                                        Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.; Arakawa, T.; Carninci, P.; Endo, T.; Fukuda, S.; Fukunishi, Y.; Hara, A.; Hayatsu, N.; Hirozane, T.; Hori, F.; Ishii, Y.; Ishikawa, J.; Ishikawa, T.; Itoh, M.; Izawa, M.; Kadota, K.; Kagawa, I.; Kai, C.; Kawai, J.; Ishikawa, J.; Kai, C.; Kawai, J.; Ishikawa, Ishika
      AUTHOR (AU):
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      TITLE (TI):
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Sciences Center(GSC), Yokohama Institute

Tel: 81-45-503-9222 Fax: 81-45-503-9216

The Institute of Physical and Chemical Research (RIKEN)

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```
Email: genome-res@gsc.riken.go.jp,
           URL:http://genome.gsc.riken.go.jp/
           Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Thermostabilization and thermoactivation of thermolabile enzymes by
           trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
             Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.
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             Automated filtration-based high-throughput plasmid preparation
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             Please visit our web site (http://genome.rtc.riken.go.jp) for
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REFERENCE:
                                                    1 (bases 1 to 316)
                                                   Konno,H.; Aizawa,K.; Akahira,S.; Akiyama,J.;
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                                                                                       Genome Science Laboratory in
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with XhoI and BamHI. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FLC I.'

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NUCLEIC ACID COUNT (NA): 102 a
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COMMENT:
       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic
       Sciences Center(GSC), Yokohama Institute
       The Institute of Physical and Chemical Research (RIKEN)
       1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsc.riken.go.jp,
       URL:http://genome.gsc.riken.go.jp/
       Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
       Thermostabilization and thermoactivation of thermolabile enzymes by
       trehalose and its application for the synthesis of full length CDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
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        High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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        Please visit our web site (http://genome.rtc.riken.go.jp) for
       further details.
REFERENCE:
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    AUTHOR (AU):
                                 Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.;
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Fukunishi,Y.; Hara,A.; Hayatsu,N.; Hirozane,T.
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Yoshida,K.; Yoshiki,A.; Yoshino,M..; Muramatsu,M.;
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Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)

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1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama. Kanagawa 230-0045, Japan
      Tel: 81-45-503-9222
      Fax: 81-45-503-9216
      Email: genome-res@gsc.riken.go.jp
      URL:http://genome.gsc.riken.go.jp/
      Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
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       Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.
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High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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Mizuno,Y.; Nakamura,M.; Oda,H.; Okazaki,Y., Ono,T.y;
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   JOURNAL (SO):
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                                                 Exploration Research Group in
                                                 Riken Genomic Sciences Center and
                                                 Genome Science Laboratory in
                                                 RIKEN. Division of Experimental
                                                 Animal Research in Riken
                                                 contributed to prepare mouse
                                                 tissues. 1st strand cDNA was primed with a primer [5'
                                                 GAGAGAGAGAGGATCCAAGAGCTCTTTTTTTT
                                                 TTTTTTVN 3'], cDNA was prepared by using trehalose
                                                 thermo-activated reverse
                                                 transcriptase and subsequently
                                                 enriched for full-length by
                                                 cap-trapper. cDNA went through one
                                                 round of subtraction to Rot =
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185.0 Second strand cDNA was prepared with the primer adapter

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L6
       ANSWER 102 OF 135
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DIVISION CODE (CI):
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DATE (DATE):
                                   23 Jun 2000
DEFINITION (DEF):
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                                  male testis Mus musculus cDNA clone 6030495A12 3
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SOURCE:
                                  house mouse.
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                                   Sciurognathi; Muridae; Murinae; Mus
NUCLEIC ACID COUNT (NA): 113 a
                                               63 c
                                                         59 q
COMMENT:
       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
       1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
       Fax: 81-45-503-9216
       Email: genome-res@gsc.riken.go.jp,
       URL:http://genome.gsc.riken.go.jp/
       Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki
       ,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
       Thermostabilization and thermoactivation of thermolabile enzymes by
       trehalose and its application for the synthesis of full length CDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
       Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki
       ,Y. and Hayashizaki,Y.
        Automated filtration-based high-throughput plasmid preparation
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        Carninci, P. and Hayashizaki, Y.
        High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
       19-44 (1999)
        Please visit our web site (http://genome.rtc.riken.go.jp) for
       further details.
REFERENCE:
                                      (bases 1 to 323)
                                 1 (bases 1 to 323)
Konno,H.; Aizawa,K.; Akahira,S.; Akiyama,J.;
Arakawa,T.; Carninci,P.; Endo,T.; Fukuda,S.;
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Mizuno,Y.; Nakamura,M.; Oda,H.; Okazaki,Y., Ono,T.y;
Owa,C.: Saito,H.: Sakai,C.: Sato,K.: Shibata,K.;
    AUTHOR (AU):
                                  Owa, C.; Saito, H.; Sakai, C.; Sato, K.; Shibata, K.; Shibata, Y.,; Shigemoto, Y.; Shinagawa, A.; Shiraki, T.;
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SEQUENCE (SEQ):

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Unpublished (2000)
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                                           Genome Science Laboratory in
                                           RIKEN. Division of Experimental
                                           Animal Research in Riken
                                           contributed to prepare mouse
                                           tissues. 1st strand cDNA was primed with a primer [5'
                                           GAGAGAGAGAGGATCCAAGAGCTCTTTTTTTT
                                           TTTTTTVN 3'], cDNA was prepared
                                           by using trehalose
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                                           transcriptase and subsequently
                                           enriched for full-length by
                                           cap-trapper. cDNA went through one
                                           round of normalization to Rot =
                                           5.0 and subtraction to Rot =
                                           100.0. Second strand cDNA was
                                           prepared with the primer adapter
of sequence [5'
                                           GAGAGAGATTCTCGAGTTAATTAAATTAATCC
                                           CCCCCCCCCC 3']. cDNA was cloned into the XhoI and BamHI sites.
                                           Vector: a modified pBluescript
                                           KS(+) after bulk excision from
                                           Lambda FLC I. Cloning sites, 5' end: SalI; 3' end: BamHI."
SEQUENCE (SEQ):
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     ANSWER 103 OF 135
                            GENBANK.RTM. COPYRIGHT 2004 on STN
LOCUS (LOC):
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GenBank ACC. NO. (GBN): BB042697
GenBank VERSION (VER):
                         BB042697.1 GI:8449083
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SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
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DIVISION CODE (CI):
                         Expressed sequence tag
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DEFINITION (DEF):
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ORGANISM (ORGN):
                         Mus musculus
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source

L6

SOURCE:

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Euteleostomi; Mammalia; Eutheria; Rodentia;
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NUCLEIC ACID COUNT (NA): 104 a
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COMMENT:
       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic
       Sciences Center(GSC), Yokohama Institute
       The Institute of Physical and Chemical Research (RIKEN)
       1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
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      Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Thermostabilization and thermoactivation of thermolabile enzymes by
      trehalose and its application for the synthesis of full length
      cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.
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        Automated filtration-based high-throughput plasmid preparation
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        Carninci, P. and Hayashizaki, Y.
        High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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        Please visit our web site (http://genome.rtc.riken.go.jp) for
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REFERENCE:
                                    (bases 1 to 289)
                                Konno,H.; Aizawa,K.; Akahira,S.; Akiyama,J.;
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    AUTHOR (AU):
                                Hori, F.; Ishii, Y.; Ishikawa, J.; Ishikawa, T.; Itoh, M.;
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                                                     RIKEN. Division of Experimental
                                                     Animal Research in Riken
                                                     contributed to prepare mouse
                                                     tissues. 1st strand cDNA was
                                                     primed with a primer [5
                                                     GAGAGAGAGGATCCAAGAGCTCTTTTTTTT
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TTTTTTTVN 3'], cDNA was prepared

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DIVISION CODE (CI):
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DATE (DATE):
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SOURCE:
                            house mouse.
 ORGANISM (ORGN):
                            Mus musculus
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NUCLEIC ACID COUNT (NA): 84 a
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                                             40 g
COMMENT:
      Contact: Yoshihide Hayashizaki
      Laboratory for Genome Exploration Research Group, RIKEN Genomic
      Sciences Center(GSC), Yokohama Institute
      The Institute of Physical and Chemical Research (RIKEN)
     1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
      Fax: 81-45-503-9216
      Email: genome-res@gsc.riken.go.jp
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     Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
      Thermostabilization and thermoactivation of thermolabile enzymes by
      trehalose and its application for the synthesis of full length
      cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.
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       High-efficiency full-length cDNA cloning, Methods Enzymol, 303,
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       Please visit our web site (http://genome.rtc.riken.go.jp) for
     further details.
REFERENCE:
                           1 (bases 1 to 237)
                           Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.; Arakawa, T.; Carninci, P.; Endo, T.; Fukuda, S.;
   AUTHOR (AU):
                           Fukunishi,Y.; Hara,A.; Hayatsu,N.; Hirozane,T.
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                                                       contributed to prepare mouse
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                                                       primed with a primer [5
                                                       GAGAGAGAAGGATCCAAGAGCTCTTTTTTTT
                                                       TTTTTTTVN 3'], cDNA was prepared by using trehalose
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into the XhoI and BamHI sites.
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                                                       Lambda FLC I. Cloning sites, 5' end: SalI; 3' end: BamHI."
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L6
      ANSWER 105 OF 135
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LOCUS (LOC):
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SEQUENCE LENGTH (SQL):
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DIVISION CODE (CI):
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DATE (DATE):
                                 19 Mar 2000
DEFINITION (DEF):
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                                 Mus musculus adaptor protein
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***Grb14*** ) mRNA, complete cds.
SOURCE:
                                 house mouse.
 ORGANISM (ORGN):
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Euteleostomi; Mammalia; Eutheria; Rodentia;
Sciurognathi; Muridae; Murinae; Mus
559 a 491 c 492 g 436 t
NUCLEIC ACID COUNT (NA): 559 a 491 c 492 REFERENCE: 1 (bases 1 to 1978)
    AUTHOR (AU):
                                 Reilly, J.F.; Mickey, G.; Maher, P.A.
                                 Association of fibroblast growth factor receptor 1 with the adaptor protein ***Grb14*** . Characterization
    TITLE (TI):
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                                 J. Biol. Chem., 275 (11), 7771-7778 (2000)
    JOURNAL (SO):
    OTHER SOURCE (OS):
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    AUTHOR (AU): TITLE (TI):
                                 Reilly, J.F.; Mickey, G.; Maher, P.A. Direct Submission
                                 Submitted (01-JUN-1999) Cell Biology, The Scripps
Research Institute, 10550 N. Torrey Pines Rd., CAL-3,
La Jolla, CA 92037, USA
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       ANSWER 106 OF 135
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COMMENT:
       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic
      Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
       Fax: 81-45-503-9216
      Email: genome-res@gsc.riken.go.jp,
      URL:http://genome.gsc.riken.go.jp/
       Sasaki,N., Izawa,M., Watahiki,M., Ozawa,K., Tanaka,T., Yoneda,Y.
      Matsuura,S., Carninci,P., Muramatsu,M., Okazaki,Y. and Hayashizaki
        Transcriptional sequencing: A method for DNA sequencing using RNA
      polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)
      Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.
        Automated filtration-based high-throughput plasmid preparation
      system. Genome Res. 9 (5), 463-470 (1999)
        Carninci, P. and Hayashizaki, Y.
        High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
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        Please visit our web site (http://genome.rtc.riken.go.jp) for
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REFERENCE:
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                               Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.; Carninci, P.; Endo, T.; Fukuda, S.; Fukunishi, Y.; Hara, A.; Hayatsu, N.; Hirozane, T.; Hori, F.; Ishii, Y.; Ishikawa, T.; Itoh, M.; Izawa, M.; Kadota, K.; Kagawa, I.; Kai, C.?; Kawai, J.; Kikuchi, N.; Kojima, Y.; Koya, S.; Kusakabe, M.; Matsuyama, T.; Miki, R.; Mizuno, Y.; Nakamura M.; Oda H.; Okazaki V.; Owa C.; Ozawa, Y.;
    AUTHOR (AU):
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    TITLE (TI):
                                RIKEN Mouse ESTs (Konno, H., et al. 1999)
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L6

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Sciences Center(GSC), Yokohama Institute

```
The Institute of Physical and Chemical Research (RIKEN)
      1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
      Fax: 81-45-503-9216
      Email: genome-res@gsc.riken.go.jp,
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      Sasaki,N., Izawa,M., Watahiki,M., Ozawa,K., Tanaka,T., Yoneda,Y.
      Matsuura,S., Carninci,P., Muramatsu,M., Okazaki,Y. and Hayashizaki
       Transcriptional sequencing: A method for DNA sequencing using RNA
      polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)
      Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki
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       Carninci, P. and Hayashizaki, Y.
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      19-44 (1999)
       Please visit our web site (http://genome.rtc.riken.go.jp) for
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REFERENCE:
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                                                   Riken Genomic Sciences Center and
                                                   Genome Science Laboratory in
                                                   RIKEN. Division of Experimental Animal Research in Riken
                                                   contributed to prepare mouse
                                                   tissues. 1st strand cDNA was primed with a primer [5'
                                                   GAGAGAGAGAGGATCCAAGAGCTCTTTTTTTT
                                                   TTTTTTTVN 3'], cDNA was prepared
                                                   by using trehalose
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                                                   enriched for full-length by
                                                   cap-trapper. cDNA went through one round of normalization to Rot = 10.0 and subtraction to Rot =
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prepared with the primer adapter

of sequence [5

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L6
      ANSWER 108 OF 135
                                  GENBANK.RTM. COPYRIGHT 2004 on STN
LOCUS (LOC):
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DATE (DATE):
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COMMENT:
       Contact: Yoshihide Hayashizaki
      Laboratory for Genome Exploration Research Group, RIKEN Genomic
      Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
      1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
      Fax: 81-45-503-9216
      Email: genome-res@gsc.riken.go.jp,
      URL:http://genome.gsc.riken.go.jp/
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      Matsuura, S., Carninci, P., Muramatsu, M., Okazaki, Y. and Hayashizaki
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       Transcriptional sequencing: A method for DNA sequencing using RNA
      polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)
      Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.
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   TITLE (TI):
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NUCLEIC ACID COUNT (NA): 105 a
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COMMENT:
     Contact: Yoshihide Hayashizaki
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Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute

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The Institute of Physical and Chemical Research (RIKEN)
      1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
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      Sasaki,N., Izawa,M., Watahiki,M., Ozawa,K., Tanaka,T., Yoneda,Y.
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                                               tissues. 1st strand cDNA was primed with a primer [5'
                                               GAGAGAGAGAGCTCTTTTTTTT
                                               TTTTTTVN 3'], cDNA was prepared by using trehalose
                                               thermo-activated reverse
                                               transcriptase and subsequently
                                               enriched for full-length by
                                               cap-trapper. cDNA went through one
                                               round of normalization to Rot =
                                               10.0 and subtraction to Rot =
                                               100.0. Second strand cDNA was
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prepared with the primer adapter of sequence [5'

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61 ttcagccata ctcttctcat tcttatacta cagaagaaga aggatgccaa aggagaatga
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181 cctgcgggtt acaaaaaaaa ataggtcaca cattgcaaat tagtgaaaac ttggattcct
     241 attacattca tgactttaaa tttattagtt aaaattaaac cttatt
L6
       ANSWER 110 OF 135
                                     GENBANK.RTM. COPYRIGHT 2004 on STN
LOCUS (LOC):
                                  AV321727
                                                     GenBank (R)
GenBank ACC. NO. (GBN): AV321727
GenBank VERSION (VER):
                                  AV321727.1 GI:6291562
CAS REGISTRY NO. (RN):
                                  247815-00-5
SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
                                  mRNA; linear
DIVISION CODE (CI):
                                  Expressed sequence tag
DATE (DATE):
                                  9 Nov 1999
DEFINITION (DEF):
                                  AV321727 RIKEN full-length enriched, 13 days embryo
                                  male_testis Mus musculus cDNA clone 6030436014 3
                                  similar to AF076619 Rattus norvegicus molecular adapter rGrb14 ( ***Grb14*** ) mRNA, mRNA sequence.
SOURCE:
                                  house mouse.
 ORGANISM (ORGN):
                                  Mus musculus
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                                  Euteleostomi; Mammalia; Eutheria; Rodentia;
                                  Sciurognathi; Muridae; Murinae; Mus
NUCLEIC ACID COUNT (NA): 84 a
                                             46 c
                                                       45 g
                                                                  78 t
COMMENT:
       Contact: Yoshihide Hayashizaki
       Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
       1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
       Fax: 81-45-503-9216
       Email: genome-res@gsc.riken.go.jp,
       URL:http://genome.gsc.riken.go.jp/
       Sasaki,N., Izawa,M., Watahiki,M., Ozawa,K., Tanaka,T., Yoneda,Y.,
       Matsuura, S., Carninci, P., Muramatsu, M., Okazaki, Y. and Hayashizaki
        Transcriptional sequencing: A method for DNA sequencing using RNA
       polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki
       ,Y. and Hayashizaki,Y.
        Automated filtration-based high-throughput plasmid preparation
       system. Genome Res. 9 (5), 463-470 (1999)
        Carninci, P. and Hayashizaki, Y.
        High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
       19-44 (1999)
        Please visit our web site (http://genome.rtc.riken.go.jp) for
       further details.
                                  1 (bases 1 to 253)
REFERENCE:
                                  Konno, H.; Aizawa, K.; Akahira, S.; Akiyama, J.;
Carninci, P.; Endo, T.; Fukuda, S.; Fukunishi, Y.; Hara, A.;
Hayatsu, N.; Hirozane, T.; Hori, F.; Ishii, Y.;
    AUTHOR (AU):
                                  Ishikawa,T.; Itoh,M.; Izawa,M.; Kadota,K.; Kagawa,I.;
Kai,C.?; Kawai,J.; Kikuchi,N.; Kojima,Y.; Koya,S.;
Kusakabe,M.; Matsuyama,T.; Miki,R.; Mizuno,Y.;
                                  Nakamura, M.; Oda, H.; Okazaki, Y.; Owa, C.; Ozawa, Y.;
                                  Saito,H.; Sano,M.; Sato,K.; Shibata,K.; Shibata,Y.; Shigemoto,Y.; Shiraki,T.; Sogabe,Y.; Sugahara,Y.;
                                 Suzuki, H.; Suzuki, H.; Takahashi, F.; Tateno, M.; Tominaga, N.; Tsunoda, Y.; Watahiki, A.; Watanabe, S.; Yamamura, T.; Yasunishi, A.; Yokota, T.; Yoshiki, A.; Yoshino, M.; Muramatsu, M.; Hayashizaki, Y. RIKEN Mouse Ests (Konno, H., et al. 1999)
    TITLE (TI):
    JOURNAL (SO):
                                  Unpublished (1999)
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SEQUENCE (SEQ):

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FEATURES (FEAT): Feature Key
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Location

Qualifier

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/lab-host="DH10B"
                                               /note="Site-1: SalI; Site-2:
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                                               and sequenced in Mouse Genome
                                               Encyclopedia Project of Genome
                                               Exploration Research Group in
                                               Riken Genomic Sciences Center and
                                               Genome Science Laboratory in
                                               RIKEN. Division of Experimental
                                               Animal Research in Riken
                                               contributed to prepare mouse
                                               tissues. 1st strand cDNA was
                                               primed with a primer [5
                                               GAGAGAGAAGGATCCAAGAGCTCTTTTTTTT
                                               TTTTTTVN 3'], cDNA was prepared
                                               by using treĥalose
                                              thermo-activated reverse
                                               transcriptase and subsequently
                                              enriched for full-length by
                                              cap-trapper. cDNA went through one
                                              round of normalization to Rot =
                                               5.0 and subtraction to Rot =
                                               100.0. Second strand cDNA was
                                              prepared with the primer adapter
                                              Vector: a modified pBluescript
                                              KS(+) after bulk excision from
                                              Lambda FLC I. Cloning sites, 5' end: Sall; 3' end: BamHI."
SEQUENCE (SEQ):
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181 tgcaaattag tgaaaacttg gattcctatt acattcatga ctttaaattt attagttaaa
   241 attaaacctt att
L6
     ANSWER 111 OF 135
                              GENBANK.RTM. COPYRIGHT 2004 on STN
LOCUS (LOC): AV259119
GenBank ACC. No. (GBN): AV259119
                                          GenBank (R)
GenBank VERSION (VER):
                           AV259119.1 GI:6246578
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SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
                           mRNA; linear
DIVISION CODE (CI):
                           Expressed sequence tag
DATE (DATE):
                           4 Nov 1999
                           AV259119 RIKEN full-length enriched, adult male testis
DEFINITION (DEF):
                            (DH10B) Mus musculus cDNA clone 4930403H14 3' similar
                           to AF076619 Rattus norvegicus molecular adapter rGrb14
                               ***Grb14*** ) mRNA, mRNA sequence.
SOURCE:
                           house mouse.
                           Mus musculus
 ORGANISM (ORGN):
                           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                           Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus
103 a 27 c 34 g 66 t
NUCLEIC ACID COUNT (NA): 103 a
COMMENT:
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Contact: Yoshihide Hayashizaki

```
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
            1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
            Tel: 81-45-503-9222
            Fax: 81-45-503-9216
            Email: genome-res@gsc.riken.go.jp,
            URL:http://genome.gsc.riken.go.jp/
            Sasaki,N., Izawa,M., Watahiki,M., Ozawa,K., Tanaka,T., Yoneda,Y.,
            Matsuura,S., Carninci,P., Muramatsu,M., Okazaki,Y. and Hayashizaki
            Transcriptional sequencing: A method for DNA sequencing using RNA polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomani Y., Canninci R., Shibata, Y., Shibata, Y
            Tomaru,Y., Carninci,P., Shibata,Y., Ozawa,Y., Muramatsu,M., Okazaki
            ,Y. and Hayashizaki,Y.
              Automated filtration-based high-throughput plasmid preparation
            system. Genome Res. 9 (5), 463-470 (1999)
              Carninci,P. and Hayashizaki,Y.
              High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
            19-44 (1999)
              Please visit our web site (http://genome.rtc.riken.go.jp) for
            further details.
                                                       1 (bases 1 to 230)
Konno,H.; Aizawa,K.; Akahira,S.; Akiyama,J.;
Carninci,P.; Endo,T.; Fukuda,S.; Fukunishi,Y.; Hara,A.;
Hayatsu,N.; Hirozane,T.; Hori,F.; Ishii,Y.;
Ishikawa,T.; Itoh,M.; Izawa,M.; Kadota,K.; Kagawa,I.;
Kai,C.?; Kawai,J.; Kikuchi,N.; Kojima,Y.; Koya,S.;
REFERENCE:
       AUTHOR (AU):
                                                        Kusakabe,M.; Matsuyama,T.; Miki,R.; Mizuno,Y.;
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                                                       Suzuki, H.; Suzuki, H.; Takahashi, F.; Tateno, M.; Tominaga, N.; Tsunoda, Y.; Watahiki, A.; Watanabe, S.; Yamamura, T.; Yasunishi, A.; Yokota, T.; Yoshiki, A.; Yoshino, M.; Muramatsu, M.; Hayashizaki, Y. RIKEN Mouse (STS (Konno, H., et al. 1999)
       TITLE (TI):
       JOURNAL (SO):
                                                        Unpublished (1999)
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                                                                                             BamHI; cDNA library was prepared
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                                                                                             Exploration Research Group in
                                                                                             Riken Genomic Sciences Center and
                                                                                             Genome Science Laboratory in
                                                                                            RIKEN. Division of Experimental Animal Research in Riken
                                                                                             contributed to prepare mouse
                                                                                             tissues. 1st strand cDNA was
                                                                                             primed with a primer [5'
                                                                                             GAGAGAGAGAGCTCTTTTTTTT
                                                                                            TTTTTTVN 3'], cDNA was prepared by using trehalose
                                                                                            thermo-activated reverse
                                                                                            transcriptase and subsequently
                                                                                            enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer
                                                                                            adapter of sequence [5
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GAGAGAGATTCTCGAGTTĀATTAAATTAATCC

CCCCCCCCC 3']. cDNA was cloned into the XhoI and BamHI sites. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FLC I. Cloning sites, 5' end: SalI; 3' end: BamHI."

ESVVEVQEAWPVGGDSRFIFRKNFAKYELFKSPP

IQNFLNAGSFPEIQGFLQLRGSGRGSGRKLWKRF

HTLFPEKMVSSCLDTPTGISHEDL

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L6
     ANSWER 112 OF 135
                           GENBANK.RTM.
                                        COPYRIGHT 2004 on STN
LOCUS (LOC):
                        AF190121
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GenBank ACC. NO. (GBN): AF190121
GenBank VERSION (VER):
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CAS REGISTRY NO. (RN):
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SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
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DIVISION CODE (CI):
                        Rodents
DATE (DATE):
                        22 Nov 2000
DEFINITION (DEF):
                        Rattus norvegicus growth factor receptor binding
                         protein GRB7 (Grb7) mRNA, complete cds.
SOURCE:
                         Norway rat.
 ORGANISM (ORGN):
                        Rattus norvegicus
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NUCLEIC ACID COUNT (NA): 440 a 602 c 548 g
                                                  462 t
                        1 (bases 1 to 2052)
REFERENCE:
   AUTHOR (AU):
                        Kasus-Jacobi, A.; Perdereau, D.; Auzan, C.; Clauser, E.;
                        Van Obberghen, E.; Mauvais-Jarvis, F.; Girard, J.;
                        Burnol,A.F.
   TITLE (TI):
                        Identification of the rat adapter ***Grb14***
                        inhibitor of insulin actions
                        J. Biol. Chem., 273 (40), 26026-26035 (1998) CA 130:20710
   JOURNAL (SO):
   OTHER SOURCE (OS):
                        2 (bases 1 to 2052)
Kasus_Jacobi,A.; Bereziat,V.; Perdereau,D.; Girard,J.;
REFERENCE:
  AUTHOR (AU):
                        Burnol, A.F.
  TITLE (TI):
                        Evidence for an interaction between the insulin
                        receptor and Grb7. A role for two of its binding
                        domains, PIR and SH2
   JOURNAL (SO):
                        Oncogene, 19 (16), 2052-2059 (2000)
  OTHER SOURCE (OS):
                        CA 133:69252
REFERENCE:
                           (bases 1 to 2052)
  AUTHOR (AU):
                        Burnol, A.F.; Perdereau, D.; Kasus-Jacobi, A.
  TITLE (TI):
                        Direct Submission
   JOURNAL (SO):
                        Submitted (27-SEP-1999) UPR 1524, CNRS, 9 rue Jules
                        Hetzel, Meudon 92190, France
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   2041 aaaaaaaaaa aa
L6
       ANSWER 113 OF 135
                                                            COPYRIGHT 2004 on STN
                                       GENBANK.RTM.
LOCUS (LOC):
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GenBank VERSION (VER):
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SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
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DIVISION CODE (CI):
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DATE (DATE):
                                    8 Mar 2000
DEFINITION (DEF):
                                    wo95a09.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone
                                    IMAGE:2463064 3' similar to TR:Q14449 Q14449
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                                                         . ;, mRNA sequence.
SOURCE:
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                                    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                                    Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                                    Hominidae; Homo
NUCLEIC ACID COUNT (NA): 90 a
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                                                          48 a
                                                                    127 t
COMMENT:
       Contact: Robert Strausberg, Ph.D.
       Email: cgapbs-r@mail.nih.gov
       Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
       Emmert-Buck, M.D., Ph.D.
         CDNA Library Preparation: M. Bento Soares, Ph.D.
        CDNA Library Arrayed by: Greg Lennon, Ph.D.
        DNA Sequencing by: Washington University Genome Sequencing Center
        Clone distribution: NCI-CGAP clone distribution information can be
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SEQUENCE (SEQ):

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Seq primer: -40UP from Gibco.
REFERENCE:
                         1 (bases 1 to 319)
   AUTHOR (AU):
                         NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
                         National Cancer Institute, Cancer Genome Anatomy
Project (CGAP), Tumor Gene Index
   TITLE (TI):
                         Unpublished (1997)
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                                          Following HAP purification, this
                                          DNA was used as tracer in a subtractive hybridization
                                          reaction. The driver was
                                          PCR-amplified cDNAs from a pool of
                                          5,000 clones made from the same
                                          library (cloneIDs 1322376-1323911, 1456007-1456775, and
                                          1500552-1502855). Subtraction by
                                          Bento Soares and M. Fatima
                                          Bonaldo.
SEQUENCE (SEQ):
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L6
     ANSWER 114 OF 135
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189 a 157 c
NUCLEIC ACID COUNT (NA): 189 a
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COMMENT:
     Contact: Robert Strausberg, Ph.D.
     Email: cgapbs-r@mail.nih.gov
     Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
     Emmert-Buck, M.D., Ph.D.
      CDNA Library Preparation: Life Technologies, Inc.
      CDNA Library Arrayed by: Greg Lennon, Ph.D.
      DNA Sequencing by: Washington University Genome Sequencing Center
      Clone distribution: NCI-CGAP clone distribution information can be
     found through the I.M.A.G.E. Consortium/LLNL at:
     www-bio.llnl.gov/bbrp/image/image.html
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    AUTHOR (AU):
                               NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
                              National Cancer Institute, Cancer Genome Anatomy
Project (CGAP), Tumor Gene Index
     TITLE (TI):
    JOURNAL (SO):
                               Unpublished (1997)
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L6
      ANSWER 115 OF 135
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                                                  COPYRIGHT 2004 on STN
LOCUS (LOC):
                              AI767914
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GenBank ACC. NO. (GBN): AI767914
GenBank VERSION (VER):
                              AI767914.1 GI:5234435
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DIVISION CODE (CI):
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DATE (DATE):
                              21 Dec 1999
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                              Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                              Hominidae; Homo
NUCLEIC ACID COUNT (NA): 112 a
                                         74 c
                                                  74 g
                                                          168 t
                                                                    1 others
COMMENT:
      Contact: Robert Strausberg, Ph.D.
      Email: cgapbs-r@mail.nih.gov
      Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
      Emmert-Buck, M.D., Ph.D.
       CDNA Library Preparation: M. Bento Soares, Ph.D.
       CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
       Clone distribution: NCI-CGAP clone distribution information can be
      found through the I.M.A.G.E. Consortium/LLNL at:
      www-bio.llnl.gov/bbrp/image/image.html
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   AUTHOR (AU):
                             NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
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TITLE (TI):
                                National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
    JOURNAL (SO):
                                Unpublished (1997)
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circles were made in vitro.
Following HAP purification, this
                                                     DNA was used as tracer in a
                                                     subtractive hybridization
                                                     reaction. The driver was PCR-amplified cDNAs from a pool of
                                                     5,000 clones made from the same
library (cloneIDs 1323912-1325831,
1471368-1472903 and
1492104-1493259). Subtraction by
                                                     Bento Soares and M. Fatima
                                                     Bonaldo.
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121 ttttcacatg ataatgttt cgcccttatt tatggtctt tattatttt cttgagtcct
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    421 agttttggg
      ANSWER 116 OF 135
L6
                                  GENBANK.RTM. COPYRIGHT 2004 on STN
LOCUS (LOC):
                               AI760945
                                                GenBank (R)
GenBank ACC. NO. (GBN): AI760945
GenBank VERSION (VER):
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DIVISION CODE (CI):
                               Expressed sequence tag
                               21 Dec 1999
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DATE (DATE):
DEFINITION (DEF):
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NUCLEIC ACID COUNT (NA): 90 a
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                                                  46 g
                                                            122 t
COMMENT:
      Contact: Robert Strausberg, Ph.D.
      Email: cgapbs-r@mail.nih.gov
      Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
      Emmert-Buck, M.D., Ph.D.
       CDNA Library Preparation: M. Bento Soares, Ph.D.
       CDNA Library Arrayed by: Greg Lennon, Ph.D.
       DNA Sequencing by: Washington University Genome Sequencing Center
       Clone distribution: NCI-CGAP clone distribution information can be
      found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html
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Std Error: 0.00

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AUTHOR (AU):
                              NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
                              National Cancer Institute, Cancer Genome Anatomy
Project (CGAP), Tumor Gene Index
    TITLE (TI):
    JOURNAL (SO):
                              Unpublished (1997)
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                                                 circles were made in vitro.
                                                 Following HAP purification, this
                                                 DNA was used as tracer in a
                                                 subtractive hybridization
                                                 reaction. The driver was PCR-amplified cDNAs from a pool of
                                                 5,000 clones made from the same
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      ANSWER 117 OF 135
L6
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LOCUS (LOC):
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                             Hominidae; Homo
NUCLEIC ACID COUNT (NA): 112 a
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COMMENT:
      Contact: Robert Strausberg, Ph.D.
      Email: cgapbs-r@mail.nih.gov
      Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
      Emmert-Buck, M.D., Ph.D.
       cDNA Library Preparation: M. Bento Soares, Ph.D.
       cDNA Library Arrayed by: Greg Lennon, Ph.D.
       DNA Sequencing by: Washington University Genome Sequencing Center
       Clone distribution: NCI-CGAP clone distribution information can be
     found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html
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      High quality sequence stop: 396.
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REFERENCE:

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap. AUTHOR (AU): National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index Unpublished (1997) TITLE (TI): JOURNAL (SO): FEATURES (FEAT): Feature Key Location Qualifier ________ /organism="Homo sapiens" /db-xref="taxon:9606" /clone="IMAGE:2296887" /clone-lib="NCIO-CGAP-Kid11" 1..408 source /lab-host="DH10B" /note="Organ: kidney; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site-1: Not I; Site-2: Eco RI; Plasmid DNA from the normalized library NCI-CGAP-Kid3 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneIDs 1322376-1323911, 1456007-1456775, and 1500552-1502855). Subtraction by Bento Soares and M. Fatima Bonaldo. SEQUENCE (SEQ): 1 tcctaaggtt taattttaac taatgaattt taaatgatga atgtaaagtc aatccaagtc 61 tttgctťátt tgcaatgcac aaacťatttt tttgtáacťt gcaggtgáaa tacattcttt 121 tcacatgata atgttttcgc ccttatttat ggtcttttat tatttttctt gagtcctttt 181 ccttcaatag tttaataagt cacttctggc ttgtctagag agcaatccta gcacaataat 241 gtttcaactt gcaaggaaga acgcccttat tgagttgata gaactccacc agctgtatta 301 gatotgtaaa tottgtgtgg coatoatoca gtgtgtggaa catttcacog toatottota 361 cccatccaca agtccttgct gaataatcaa tcgctgagcc tcatctta COPYRIGHT 2004 on STN L6 ANSWER 118 OF 135 GENBANK.RTM. LOCUS (LOC): AI671320 GenBank (R) GenBank ACC. NO. (GBN): AI671320 GenBank VERSION (VER): AI671320.1 GI:4851051 CAS REGISTRY NO. (RN): 232699-27-3 SEQUENCE LENGTH (SQL): 497 MOLECULE TYPE (CI): mRNA; linear DIVISION CODE (CI): Expressed sequence tag 17 Dec 1999
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. mpNA sequence. DATE (DATE): DEFINITION (DEF): SOURCE: human. ORGANISM (ORGN): Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo NUCLEIC ACID COUNT (NA): 144 a 100 c 87 g 164 t 2 others COMMENT: Contact: Robert Strausberg, Ph.D. Email: cgapbs-r@mail.nih.gov Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D. cDNA Library Preparation: M. Bento Soares, Ph.D. cDNA Library Arrayed by: Greg Lennon, Ph.D. DNA Sequencing by: Washington University Genome Sequencing Center Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html Insert Length: 795 Std Error: 0.00 Seq primer: -40UP from Gibco High quality sequence stop: 372. REFERENCE: 1 (bases 1 to 497)

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

AUTHOR (AU):

National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index TITLE (TI): Unpublished (1997) JOURNAL (SO): FEATURES (FEAT): Location Qualifier Feature Key _________________________________ /organism="Homo sapiens" /db-xref="taxon:9606" /clone="IMAGE:2316554" /clone-lib="NCI-CGAP-Kid11" /lab-host="DH10B" source 1..497 /note="Organ: kidney; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site-1: Not I; Site-2: Eco RI; Plasmid DNA from the normalized library NCI-CGAP-Kid3 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5.000 clones made from the same library (cloneIDs 1322376-1323911, 1456007-1456775, and 1500552-1502855). Subtraction by Bento Soares and M. Fatima Bonaldo. SEQUENCE (SEQ): 1 gcctgccagt gacacataaa tatcactatt gccaaattcg ctaaaaactg caaatgccgc 61 ggttcctttg atgttccttt agtagaaaaa tataaaccag atcttcttag aaaaaagtaa 121 atttttttcc aagacttctt teeetgttet ttegeatgta agaaaccatg aattteagga 181 tatgtgcttg aactcagaaa catctgcaaa atctgtgtgg gggatatttc accattggtt 241 tcagttgcaa aagataccat atgctctgga aaaaaataca ttgggttttt aaagaactca 301 tatitggcat aaitttttct aaagtatagt ttgttttctt cttctatccc ccagttggat 361 agcacticaa tcaccagttc gtggtcttct atigttcttt ctacacctat gtgaggcagg 421 tgctcanaaa gggtccagct gtggtcatca atgtaatgat tcttcaggat caacagctga 481 ncaacatctc gagccgt ANSWER 119 OF 135 GENBANK.RTM. COPYRIGHT 2004 on STN L6 GenBank (R) LOCUS (LOC): AI624682 GenBank ACC. NO. (GBN): AI624682 GenBank VERSION (VER): AI624682.1 GI:4649613 230649-94-2 CAS REGISTRY NO. (RN): SEQUENCE LENGTH (SQL): 533 MOLECULE TYPE (CI): DIVISION CODE (CI): mRNA; linear Expressed sequence tag 14 Dec 1999 DATE (DATE): ts43e12.x1 NCI_CGAP_Ut1 Homo sapiens cDNA clone DEFINITION (DEF): IMAGE:2231374 3' similar to TR:Q14449 Q14449 ***GRB14*** . ;, mRNA sequence. SOURCE: ORGANISM (ORGN): Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Hómo 144 a 104 c NUCLEIC ACID COUNT (NA): 144 a 91 g **1**93 t 1 others COMMENT: Contact: Robert Strausberg, Ph.D. Email: cgapbs-r@mail.nih.gov Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D. cDNA Library Preparation: Life Technologies, Inc.

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                               NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
                              National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
     TITLE (TI):
     JOURNAL (SO):
                              Unpublished (1997)
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      ANSWER 120 OF 135
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LOCUS (LOC):
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DIVISION CODE (CI):
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DATE (DATE):
                              13 May 1999
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SOURCE:
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                              Hominidae; Homo
NUCLEIC ACID COUNT (NA): 179 a
                                         144 c
                                                  132 g
                                                            242 t
                                                                      4 others
COMMENT:
      Contact: Robert Strausberg, Ph.D.
      Email: cgapbs-r@mail.nih.gov
      Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
      Emmert-Buck, M.D., Ph.D.
       CDNA Library Preparation: Life Technologies, Inc.
       CDNA Library Arrayed by: Greg Lennon, Ph.D.
       DNA Sequencing by: Washington University Genome Sequencing Center
      Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html
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REFERENCE:
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   AUTHOR (AU):
                             NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
   TITLE (TI):
                             National Cancer Institute, Cancer Genome Anatomy
                             Project (CGAP), Tumor Gene Index
   JOURNAL (SO):
                             Unpublished (1997)
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 L6
       ANSWER 121 OF 135
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 LOCUS (LOC):
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 GenBank ACC. NO. (GBN): AI609572
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MOLECULE TYPE (CI):
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DATE (DATE):
                                 14 May 1999
                                 tw28a08.x1 NCI_CGAP_Ov35 Homo sapiens cDNA clone
DEFINITION (DEF):
                                 IMAGE:2260982 3' similar to TR:Q14449 Q14449
                                    ***GRB14*** ;, mRNA sequence.
SOURCE:
                                 human.
 ORGANISM (ORGN):
                                Homo sapiens
                                Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                                Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                                Hominidae; Homo
NUCLEIC ACID COUNT (NA): 100 a
                                                      57 g
                                                               151 t
                                                                          1 others
COMMENT:
       Contact: Robert Strausberg, Ph.D.
       Email: cgapbs-r@mail.nih.gov
       Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
       R. Emmert-Buck, M.D., Ph.D.
        CDNA Library Preparation: Life Technologies, Inc.
        CDNA Library Arrayed by: Greg Lennon, Ph.D.
        DNA Sequencing by: Washington University Genome Sequencing Center
        Clone distribution: NCI-CGAP clone distribution information can be
      found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html
       Insert Length: 318
                                 Std Error: 0.00
       Seq primer: -40UP from Gibco
      High quality sequence stop: 324
      POLYA=No.
REFERENCE:
                                1 (bases 1 to 368)
    AUTHOR (AU):
                                NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
    TITLE (TI):
                                National Cancer Institute, Cancer Genome Anatomy
                                Project (CGAP), Tumor Gene Index
    JOURNAL (SO):
                                Unpublished (1997)
FEATURES (FEAT):
  Feature Key
                          Location
                                                          Qualifier
source
                     1..368
                                                     /organism="Homo sapiens"
/db-xref="taxon:9606"
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/clone="IMAGE:2260982"
                                            /clone-lib="NCI-CGAP-0v35"
                                            /tissue-type="tumor, 5 pooled (see
                                            description)"
                                            /lab-host="DH10B"
                                            /note="Organ: ovary; Vector:
pCMV-SPORT6; Site-1: SalI; Site-2:
NotI; This library represents the
                                            normalized version of
                                            NCI-CGAP-Ov23. Cloned unidirectionally. Primer: Oligo
                                            dT. Average insert size 0.86 kb.
                                            Tumor types include: mixed
                                            Mullerian tumor, papillary serous,
                                            clear cell, spindle cell. All are
                                            primary tumors, metastasis
                                            positive. Constructed by Life
                                            Technologies.
     1 tttttttttt ttttttttt ttttttatgc atacacttct tggatttatt aatgctatag
    61 ttctatgaaa tccatgagta aatatagaaa cattgaaatt ccttctctct ctttagagtt
   121 ttcttggtac gggatágtca gagtaacccc aaaactttcg tactgtcaat gagtcátgga
   181 caaaaaataa agcactttca aattatacca gtaagtaatt cgtgatttca catttgtgta
   241 tragaaatga ccttaatgct aagcttttga tcttaatgca taagcttttg gaaactttgg
   301 ttttcttttg gnctttttat taaatataat ttggcagctt gtgctttgac tagagccccg
                            GENBANK.RTM. COPYRIGHT 2004 on STN
                          AI522272
                                         GenBank (R)
GenBank ACC. NO. (GBN): AI522272
                          AI522272.1 GI:4436407
                          228602-27-5
                          mRNA; linear
                          Expressed sequence tag
                          13 Apr 1999
                          ti84g01.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:2138736 3' similar to TR:Q14449 Q14449 ***GRB14*** .;, mRNA sequence.
                          Homo sapiens
                          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                          Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                          Hominidae; Homo
                                    118 c
                                             118 a
                                                      208 t
     Contact: Robert Strausberg, Ph.D.
     Email: cgapbs-r@mail.nih.gov
     Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
      cDNA Library Preparation: M. Bento Soares, Ph.D.
      CDNA Library Arrayed by: Greg Lennon, Ph.D.
      DNA Sequencing by: Washington University Genome Sequencing Center
      Clone distribution: NCI-CGAP clone distribution information can be
     found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html Insert Length: 1380 Std Error: 0.00
     Seq primer: -40UP from Gibco
     High quality sequence stop: 308.
                          1 (bases 1 to 604)
                          NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
                          National Cancer Institute, Cancer Genome Anatomy
                          Project (CGAP), Tumor Gene Index
                          Unpublished (1997)
                                                Qualifier
/organism="Homo sapiens"
/db-xref="taxon:9606"
                                            /clone="IMAGE:2138736"
                                            /clone-lib="NCI-CGAP-Kid11"
                                            /lab-host="DH10B"
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/note="Organ: kidney; Vector:

SEQUENCE (SEQ):

361 cgtccgcc

GenBank VERSION (VER):

CAS REGISTRY NO. (RN):

SEQUENCE LENGTH (SQL):

MOLECULE TYPE (CI):

DIVISION CODE (CI):

DEFINITION (DEF):

ORGANISM (ORGN):

LOCUS (LOC):

DATE (DATE):

SOURCE:

COMMENT:

REFERENCE:

source

AUTHOR (AU):

JOURNAL (SO):

TITLE (TI):

FEATURES (FEAT):

Feature Key

ANSWER 122 OF 135

NUCLEIC ACID COUNT (NA): 160 a

Emmert-Buck, M.D., Ph.D.

Location

1..604

604

L6

```
pT7T3D-Pac (Pharmacia) with a modified polylinker; Site-1: Not I; Site-2: Eco RI; Plasmid DNA from the normalized library NCI-CGAP-Kid3 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneIDs 1322376-1323911, 1456007-1456775, and 1500552-1502855). Subtraction by Bento Soares and M. Fatima Bonaldo. "
```

SEQUENCE (SEQ): 1 ttcctaaggt ttaattttaa ctaatgaatt ttaaatgatg aatgtaaagt caatccaagt 61 ctttgcttat ttgcaatgca caaactattt ttttgtaact tgcaggtgaa atacattctt 121 ttcacatgat aatgttticg cccttattta tggtctttta tiatiitict tgagtccttt 181 tectteaata gtttaataag teaettetgg ettgtetaga gageaateet ageacaataa 241 tgtttcaact tgcaaggaag aacgccctta ttgagttgat agaactccac cagctgtatt 301 agatetgtaa atettgtgtg gecateatee agtgtgtgga acattteace gteatettet 361 actggtataa tttgaaagtg ctttattttt tgtccatgac tcattgacag tacgaaagtt 421 ttggggttac tctgactatc ccgtaccaag aaaactccat ccacaagtcc ttgctgaata 481 atcaatcgct gagcctcatc tctagaaatt ttgtggtgaa accatggctg ggacccgtgg 541 atagccatgt ttgtggcaaa gcttctggaa gaggcagggg ggctccggga gtgcccaggc 601 qtaa L6 ANSWER 123 OF 135 GENBANK.RTM. COPYRIGHT 2004 on STN LOCUS (LOC): AI505286 GenBank (R) GenBank ACC. NO. (GBN): AI505286 GenBank VERSION (VER): AI505286.1 GI:4403137 CAS REGISTRY NO. (RN): 228210-72-8 SEQUENCE LENGTH (SQL): MOLECULE TYPE (CI): DIVISION CODE (CI): 578 mRNA; linear Expressed sequence tag 11 Mar 1999 DATE (DATE): vp98h08.x1 Stratagene mouse diaphragm (#937303) Mus **DEFINITION (DEF):** musculus cDNA clone IMAGE:1092831 3 similar to TR:Q14449 Q14449 ***GRB14*** . ;, mRNA sequence. house mouse. SOURCE: ORGANISM (ORGN): Mus musculus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus 148 a 135 c 111 g 179 t 5 others NUCLEIC ACID COUNT (NA): 148 a COMMENT: Contact: Marra M/WashU-NCI Mouse EST Project 1999 Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800 Fax: 314 286 1810 Email: mouseest@watson.wustl.edu This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information. MGI:599063 This clone was previously sequenced on the 5' end only, this new data is from the 3' end High quality sequence stop: 379. REFERENCE: 1 (bases 1 to 578) AUTHOR (AU): Marra, M.; Hillier, L.; Kucaba, T.; Martin, J.; Beck, C.; Wylie, T.; Underwood, K.; Steptoe, M.; Theising, B.; Allen,M.; Bowers,Y.; Person,B.; Swaller,T.; Gibbons,M.; Pape,D.; Harvey,N.; Schurk,R.; Ritter,E.; Kohn,S.; Shin,T.; Jackson,Y.; Cardenas,M.; McCann,R.; Waterston, R.; Wilson, R. The WashU-NCI Mouse EST Project 1999 TITLE (TI): JOURNAL (SO): Unpublished (1999)

Qualifier

FEATURES (FEAT):

Feature Key

Location

source

1..368

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/organism="Mus musculus"
/db-xref="taxon:10090"
/clone="IMAGE:1092831"
/clone-lib="Stratagene mouse
diaphragm (#937303)
/tissue-type="diaphragm"
/dev-stage="adult
/lab-host="SOLR (kanamycin
resistant)"
/note="Organ: diaphragm; Vector:
pBluescript SK-; Site-1: EcoRI;
Site-2: XhoI; Cloned unidirectionally from mRNA
prepared from diaphragm muscle.
Primer: Oligo dT. Average insert
size: 1.5 kb. Uni-ZAP XR Vector;
~5' adaptor sequence: 5'
GAATTCGGCACGAG 3' ~3' adaptor
sequence: 5
```

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SEQUENCE (SEQ):
   1 aataaggttt aattttaact aataaattta aaggcatgag tgtaatagga atccaagttt 61 tcactaattt gcaatgtgtg acctattttt ttttgtaacc cgcaggtgaa atcttctttt 121 caccatggtt tgttttcagc cttgtgatct ctctctct ctctaatcat tctcctttgc
   181 atccttette tietgtagig taacgagtga cacacagttt ggetaaacag ccatectage
   241 acagtaatgo ttcagottgo aaggaaggac occootgttg agotggtaga actocaccag
   301 ctggatgagg tetgigaact tegiatggee ateateeaga giatggaaca geteaceate
   361 atčítcťačď ggtataatnt gaťagtďítt tatcttttďt čcatďáctca ítgacagtac
   421 aaaagttctg gggttactct gactatcccg taccaagaaa actncatcca caggcccctg
481 ccgaatgatc agncgctgag cctcatctct tgaaatnctg tggtgaaacc catgttggac
   541 cgatggaaac catgttcaca ccanaactct ggaagggc
     ANSWER 124 OF 135
                                              COPYRIGHT 2004 on STN
L6
                              GENBANK.RTM.
LOCUS (LOC):
                            AI494168
                                            GenBank (R)
GenBank ACC. NO. (GBN): AI494168
GenBank VERSION (VER):
                            AI494168.1 GI:4395171
CAS REGISTRY NO. (RN):
                            228132-01-2
SEQUENCE LENGTH (SQL):
                            368
MOLECULE TYPE (CI):
DIVISION CODE (CI):
                            mRNA; linear
                            Expressed sequence tag
                            13 Apr 1999
DATE (DATE):
DEFINITION (DEF):
                            ti14f01.y1 NCI_CGAP_Kid11 Homo sapiens cDNA clone
                            IMAGE:2130457 5' similar to TR:Q14449 Q14449 ***GRB14*** ;, mRNA sequence.
SOURCE:
                            human.
 ORGANISM (ORGN):
                            Homo sapiens
                            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                            Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                            Hominidae; Homo
NUCLEIC ACID COUNT (NA): 98 a
                                      71 c
                                              58 q
                                                      141 t
COMMENT:
      Contact: Robert Strausberg, Ph.D.
      Email: cgapbs-r@mail.nih.gov
      Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
      Emmert-Buck, M.D., Ph.D.
       CDNA Library Preparation: M. Bento Soares, Ph.D.
       CDNA Library Arrayed by: Greg Lennon, Ph.D.
       DNA Sequencing by: Washington University Genome Sequencing Center
     Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html
      Insert Length: 416
                              Std Error: 0.00
      Seq primer: -40RP from Gibco.
REFERENCE:
                                (bases 1 to 368)
   AUTHOR (AU):
                            NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
   TITLE (TI):
                            National Cancer Institute, Cancer Genome Anatomy
                            Project (CGAP), Tumor Gene Index
   JOURNAL (SO):
                            Unpublished (1997)
FEATURES (FEAT):
  Feature Key
                       Location
                                                    Qualifier
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/organism="Homo sapiens"

/db-xref="taxon:9606'

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/clone="IMAGE:2130457"
                                          /clone-lib="NCI-CGAP-Kid11"
/lab-host="DH10B"
                                          /note="organ: kidney; Vector:
                                          pT7T3D-Pac (Pharmacia) with a
                                          modified polylinker; Site-1: Not
                                          I; Site-2: Eco RI; Plasmid DNA
                                          from the normalized library
                                          NCI-CGAP-Kid3 was prepared, and ss circles were made in vitro.
                                          Following HAP purification, this
                                          DNA was used as tracer in a subtractive hybridization
                                          reaction. The driver was
                                          PCR-amplified cDNAs from a pool of
                                          5,000 clones made from the same
                                          library (cloneIDs 1322376-1323911, 1456007-1456775, and
                                          1500552-1502855). Subtraction by
                                          Bento Soares and M. Fatima
                                          Bonaldo.
  1 tcctaaggtt taattttaac taatgaattt taaatgatga atgtaaagtc aatccaagtc
 61 tttgcttatt tgcaatgcac aaactattt tttgtaactt gcaggtgaaa tacattcttt
121 tcacatgata atgitticge eettatitat ggiettitat tattitiett gagieettit
181 ccttcaatag tttaataagt cacttctggc ttgtctagag agcaatccta gcacaataat
241 gtttcaactt gcaaggaaga acgcccttat tgagttgata gaactccacc agctgtatta
301 gatctgtaaa tcttgtgtgg ccatcatcca gtgtgtggaa catttcaccg tcatcttctc
                          GENBANK.RTM. COPYRIGHT 2004 on STN
                                      GenBank (R)
                                     GI:4271348
                        Expressed sequence tag
                       15 Mar 2000
my18a09.y1 Barstead mouse heart MPLRB3 Mus musculus
cDNA clone IMAGE:696184 5' similar to TR:Q14449 Q14449
                          ***GRB14*** . ;, mRNA sequence.
                        Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                        Euteleostomi; Mammalia; Eutheria; Rodentia;
                       Sciurognathi; Muridae; Murinae; Mus
                                           143 g
                                 171 c
  Contact: Marra M/WashU-NCI Mouse EST Project 1999
  Washington University School of Medicine
  4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
  Email: mouseest@watson.wustl.edu
  This clone is available royalty-free through LLNL; contact the
  IMAGE Consortium (info@image.llnl.gov) for further information.
  This read is a RESEQUENCE of a previously sequenced mouse clone
  This read has been verified (found to hit its original self in the
                       1 (bases 1 to 503)
                       Marra,M.; Hillier,L.; Kucaba,T.; Martin,J.; Beck,C.;
                       Wylie, T.; Underwood, K.; Steptoe, M.; Theising, B.;
                       Allen,M.; Bowers,Y.; Person,B.; Swaller,T.;
Gibbons,M.; Pape,D.; Harvey,N.; Schurk,R.; Ritter,E.;
Kohn,S.; Shin,T.; Jackson,Y.; Cardenas,M.; McCann,R.;
                       Waterston, R.; Wilson, R.
                       The WashU-NCI Mouse EST Project 1999
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SEQUENCE (SEQ):

361 ctcgtgcc

LOCUS (LOC):

DATE (DATE): **DEFINITION (DEF):**

SOURCE:

COMMENT:

ANSWER 125 OF 135

GenBank VERSION (VER):

CAS REGISTRY NO. (RN): SEQUENCE LENGTH (SQL):

MOLECULE TYPE (CI):

DIVISION CODE (CI):

ORGANISM (ORGN):

GenBank ACC. NO. (GBN): AI425417

NUCLEIC ACID COUNT (NA): 103 a

Tel: 314 286 1800 Fax: 314 286 1810

correct orientation)

Seq primer: -40RP from Gibco High quality sequence stop: 493

MGI:429744

POLYA=No.

AUTHOR (AU):

TITLE (TI): JOURNAL (SO):

REFERENCE:

AI425417

503

AI425417.1

226504-01-4

mRNA; linear

house mouse.

Mus musculus

Unpublished (1999)

L6

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FEATURES (FEAT):
 Feature Key
                    Location
                                            Qualifier
/organism="Mus musculus"
source
                                        /strain="BALB/c"
/db-xref="taxon:10090"
                                        /clone="IMAGE:696184"
                                         /clone-lib="Barstead mouse heart
                                        MPLRB3
                                        /sex="mixed"
                                        /tissue-type="heart"
/dev-stage="6 weeks"
                                         /lab-host="DH10B"
                                         /note="Organ: heart; Vector:
                                        pT7T3D-Pač (Pharmacia) with a
                                        modified polylinker; Site-1: EcoRI; Site-2: NotI; 1st strand
                                        cDNA was primed with a Not I -
                                        oligo(dT) primer [5
                                        TGTTACGAATCTGAAGTGGGAGCGGCCGCCCTTT
                                        to Eco RI adaptors
                                        [CTTGGATTCGGTACC], digested with Not I and cloned into the Not I
                                        and Eco RI sites of the modified
                                        pT7T3 vector. Library constructed
                                        by Bob Barstead.
SEQUENCE (SEQ):
     1 aatteggate caaggeaagg egetegetge etgeaacege teggetetge tegeeeceag
  301 ggagaaaaat gaaagatctg gatgttctgg aaaagccacc cattcccaac ccctttcctg
   361 agctetgetg eteteegett acatetgtge tgteageagg cetgtttece agggecaatt
   421 caaggaagaa gcaggtgatt aaagtttaca gcgaggatga aaccagcaga gcattagagg
   481 tgcccagtga catcacagcc cga
L6
     ANSWER 126 OF 135
                          GENBANK.RTM. COPYRIGHT 2004 on STN
                        AI383743
LOCUS (LOC):
                                     GenBank (R)
GenBank ACC. NO. (GBN): AI383743
GenBank VERSION (VER):
                        AI383743.1 GI:4196524
CAS REGISTRY NO. (RN):
                        225333-52-8
SEQUENCE LENGTH (SQL):
                        423
MOLECULE TYPE (CI):
                        mRNA; linear
DIVISION CODE (CI):
                        Expressed sequence tag
                        18 Mar 1999
DATE (DATE):
                        tc47e05.x1 Soares_total_fetus_Nb2HF8_9w Homo sapiens cDNA clone IMAGE:2067776 3' similar to TR:Q14449 Q14449
DEFINITION (DEF):
                          ***GRB14*** . ;, mRNA sequence.
SOURCE:
                        human.
                        Homo sapiens
ORGANISM (ORGN):
                        Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                        Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                        Hominidae; Homo
116 a 74 c
NUCLEIC ACID COUNT (NA): 116 a
                                        69 g
                                               162 t 2 others
COMMENT:
     Contact: Robert Strausberg, Ph.D.
     Email: cgapbs-r@mail.nih.gov
     This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information. Insert Length: 967 Std Error: 0.00
     Seg primer: -40UP from Gibco.
REFERENCE:
                        1 (bases 1 to 423)
  AUTHOR (AU):
                        NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
   TITLE (TI);
                        National Cancer Institute, Cancer Genome Anatomy
                        Project (CGAP), Tumor Gene Index
   JOURNAL (SO):
                        Unpublished (1997)
FEATURES (FEAT):
 Feature Key Location
                                            Qualifier
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/organism="Homo sapiens"

source 1..423

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/clone="IMAGE:2067776"
                                           /clone-lib="Soares-total-fetus-Nb2
                                           HF8-9w"
                                           /dev-stage="8-9 weeks"
                                           /lab-hosť="DH10B"
                                           /note="Vector: pT7T3D-Pac
                                           (Pharmacia) with a modified
                                           polylinker; Site-1: Not I; Site-2:
                                           Eco RI; 1st strand cDNA was
                                           prepared from mRNA obtained from
                                           pooled 8-9 week (total) fetus
                                           material with a Not I - oligo(dT)
                                           primer [5
                                           TGTTACCĀATCTGAAGTGGGAGCGGCCGCTTAAT
                                           זוודדדדדדדדד 3'].
                                           Double-stranded cDNA was ligated
                                           to Eco RI adaptors (Pharmacia),
                                           digested with Not I and cloned
                                           into the Not I and Eco RI sites of
                                           the modified pT7T3 vector. Library
                                           went through one round of
                                           normalization, and was constructed
                                            by Bento Soares and M. Fatima
                                           Bonaldo.
  1 cctaaggttt aattttaact aatgaatttt aaatgatgaa tgtaaagtca atccaagtct
 61 ttgcttättt gcaatgcaca aactattttt ttgtäacttg cäggtgäaat acattctttt
121 cacatgataa cgttttcgcc cttatttatg gtcttttatt attittcttg agtccttttc 181 cttcaatagt ttaataagtc acttctggct tgtctagaga gcaatcctag cacaataatg
241 tttcaactig caaggaagaa cgccctiatt gagttgatag aactccacca gctgtattag 301 atctgtaaat cttgtgtggc catcatccag tgtgtggaac atttcaccgt catcttctac
361 tggtataatt ngaaagtgct ttattntttt gtcatgactc attgacagta caaaagtttt
                          GENBANK RTM. COPYRIGHT 2004 on STN
                                       GenBank (R)
                        AI364971.1 GI:4124660
                        Expressed sequence tag
                        qz41h03.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone
                        IMAGE:2029493 3' similar to TR:Q14449 Q14449
                          ***GRB14*** . ;, mRNA sequence.
                        Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                        Euteleostomi; Mammalia; Euthéria; Primates; Catarrhini;
                        Hominidae; Homo
                                         48 a
                                                 125 t
                                                           1 others
  Contact: Robert Strausberg, Ph.D.
  Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
   cDNA Library Preparation: M. Bento Soares, Ph.D. cDNA Library Arrayed by: Greg Lennon, Ph.D. DNA Sequencing by: Washington University Genome Sequencing Center
   Clone distribution: NCI-CGAP clone distribution information can be
  found through the I.M.A.G.E. Consortium/LLNL at:
  www-bio.llnl.gov/bbrp/image/image.html
                          Std Error: 0.00
                           (bases 1 to 318)
                        NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
                        National Cancer Institute, Cancer Genome Anatomy
Project (CGAP), Tumor Gene Index
                        Unpublished (1997)
```

/db-xref="taxon:9606"

SEQUENCE (SEQ):

421 ggg

LOCUS (LOC):

DATE (DATE):

SOURCE:

COMMENT:

REFERENCE:

AUTHOR (AU):

TITLE (TI):

FEATURES (FEAT):

Feature Key

JOURNAL (SO):

ANSWER 127 OF 135

GenBank VERSION (VER):

CAS REGISTRY NO. (RN):

SEQUENCE LENGTH (SQL):

MOLECULE TYPE (CI):

DIVISION CODE (CI):

DEFINITION (DEF):

ORGANISM (ORGN):

GenBank ACC. NO. (GBN): AI364971

NUCLEIC ACID COUNT (NA): 90 a

Insert Length: 447

Email: cgapbs-r@mail.nih.gov

Seq primer: -40UP from Gibco.

Location

Emmert-Buck, M.D., Ph.D.

AI364971

318

human.

224494-55-7

mRNA; linear

16 Feb 1999

Homo sapiens

54 c

Qualifier

L6

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/db-xref="taxon:9606"
/clone="IMAGE:2029493"
/clone-lib="NCI-CGAP-Kid11"
                 1..318
source
                                            /lab-host="DH10B"
                                            /note="Organ: kidney; Vector:
                                           pT7T3D-Pac (Pharmacia) with a
                                           modified polylinker; Site-1: Not
                                           I; Site-2: Eco RI; Plasmid DNA
                                           from the normalized library
                                           NCI-CGAP-Kid3 was prepared, and ss circles were made in vitro.
                                           Following HAP purification, this DNA was used as tracer in a
                                           subtractive hybridization
                                           reaction. The driver was
                                           PCR-amplified cDNAs from a pool of
                                           5,000 clones made from the same
                                           library (cloneIDs 1322376-1323911, 1456007-1456775, and
                                           1500552-1502855). Subtraction by
                                           Bento Soares and M. Fatima
                                           Bonaldo.
SEQUENCE (SEQ):
     1 ttcctaaggt ttaattttaa ctaatgaatt ttaaatgatg aatgtaaagt caatccaagt
    61 ctttgcttat ttgcaatgca caaactattt ttttgtaact tgcaggtgaa atacattctt
   121 ttcacatgat aatgitticg cccttatita tggnctitta ttattitict tgagicctit
   181 teetteaata gittaataag teaetteigg etigtetaga gageaateet ageacaataa
   241 tgtttcaact tgcaaggaag aacgccctta ttgagttgat agaactccac cagctgtatt
   301 agatctgtaa atcttgtg
     ANSWER 128 OF 135
                            GENBANK.RTM. COPYRIGHT 2004 on STN
L6
LOCUS (LOC):
                          AI263214
                                        GenBank (R)
GenBank ACC. NO. (GBN): AI263214
GenBank VERSION (VER):
                          AI263214.1 GI:3871417
CAS REGISTRY NO. (RN):
                          221598-25-0
SEQUENCE LENGTH (SQL):
                          382
MOLECULE TYPE (CI):
                          mRNA; linear
DIVISION CODE (CI):
                          Expressed sequence tag
                          3 Feb 1999
DATE (DATE):
DEFINITION (DEF):
                          qz36f04.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone
                          IMAGE:2028991 3' similar to TR:Q14449 Q14449
                            ***GRB14*** . ;, mRNA sequence.
SOURCE:
                          human.
                          Homo sapiens
 ORGANISM (ORGN):
                          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                          Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                          Hominidae; Homo
NUCLEIC ACID COUNT (NA): 112 a
                                           56 g
                                                   154 t
COMMENT:
     Contact: Robert Strausberg, Ph.D.
     Email: cgapbs-r@mail.nih.gov
     Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
     Emmert-Buck, M.D., Ph.D.
      CDNA Library Preparation: M. Bento Soares, Ph.D.
      CDNA Library Arrayed by: Greg Lennon, Ph.D.
      DNA Sequencing by: Washington University Genome Sequencing Center
     Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html
     Insert Length: 730
                           Std Error: 0.00
     Seq primer: -40UP from Gibco
     High quality sequence stop: 381.
REFERENCE:
                          1 (bases 1 to 382)
   AUTHOR (AU):
                          NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
   TITLE (TI):
                          National Cancer Institute, Cancer Genome Anatomy
                          Project (CGAP), Tumor Gene Index
   JOURNAL (SO):
                          Unpublished (1997)
FEATURES (FEAT):
  Feature Key
                     Location
                                               Qualifier
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/organism="Homo sapiens"

1..382

source

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/db-xref="taxon:9606"
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/clone-lib="NCI-CGAP-Kid11"
/lab-host="DH10B"
                                             /note="organ: kidney; Vector:
                                             pT7T3D-Pac (Pharmacia) with a
                                             modified polylinker; Site-1: Not
                                             I; Site-2: Eco RI; Plasmid DNA
                                             from the normalized library
                                            NCI-CGAP-Kid3 was prepared, and ss
circles were made in vitro.
                                            Following HAP purification, this DNA was used as tracer in a
                                             subtractive hybridization
                                             reaction. The driver was
                                             PCR-amplified cDNAs from a pool of
                                             5,000 clones made from the same
                                             library (cloneIDs 1322376-1323911, 1456007-1456775, and
                                             1500552-1502855). Subtraction by
                                             Bento Soares and M. Fatima
                                             Bonaldo.
     1 tttttttttt ttttttttt cctaaggttt aattttaact aatgaatttt aaatgatgaa
    61 tgtaaagtca atccaagtct ttgcttattt gcaatgcaca aactattttt ttgtaacttg
   121 caggggaaat acattetttt cacatgataa igttticgee ettatttatq gicttttati
   181 attiticttg agtccttttc cttcaatagt tiaataagtc acttctggci igtctagaga
   241 gcaatcctag cacaataatg tttcaacttg caaggaaaaa cgcccttatt gagttgatag 301 aactcacaca cacgcccctt tggggtttta attttttaaa aggaaaaatt tcccggttgg
                            GENBANK.RTM. COPYRIGHT 2004 on STN
                                         GenBank (R)
                          AF076619.1 GI:3650499
                          mRNA; linear
                          Rattus norvegicus molecular adapter rGrb14 (
                             ***Grb14*** ) mRNA, complete cds.
                          Rattus norvegicus
                          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                          Euteleostomi; Mammalia; Eutheria; Rodentia;
                          Sciurognathi; Muridae; Murinae; Rattus
                                   460 c
                                             500 g
                                                      444 t
                          1 (bases 1 to 1950)

Kasus-Jacobi,A.; Perdereau,D.; Auzan,C.; Clauser,E.;

Van Obberghen,E.; Mauvais-Jarvis,F.; Girard,J.;
                          Identification of the rat adapter ***Grb14***
                                                                                   as an
                          inhibitor of insulin actions
                          J. Biol. Chem., 273 (40), 26026-26035 (1998)
                          CA 130:20710
                              (bases 1 to 1950)
                          Kasus-Jacobi, A.; Perdereau, D.; Burnol, A.-F.
                          Direct Submission
                          Submitted (03-JUL-1998) UPR 1524, CNRS, 9 rue Jules
                          Hetzel, Meudon 92190, France
                                                 Qualifier
/organism="Rattus norvegicus"
                                            /strain="Wistar"
/db-xref="taxon:10116"
                                            /gene="Grb14"
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                                            /note="signal transduction
                                            protein; Grb7 family member; binds
                                            the insulin receptor"
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/codon-start=1

SEQUENCE (SEQ):

LOCUS (LOC):

DATE (DATE): DEFINITION (DEF):

SOURCE:

REFERENCE:

REFERENCE:

source

gene

CDS

L6

361 gggttttnaa aaaaaaaaaa aa

GenBank ACC, NO. (GBN): AF076619

NUCLEIC ACID COUNT (NA): 546 a

AF076619

1950

Rodents 26 Sep 1998

216295-93-1

Norway rat.

Burnol, A.F.

Location

1..1950

1..1950

70..1686

ANSWER 129 OF 135

GenBank VERSION (VER):

CAS REGISTRY NO. (RN):

SEQUENCE LENGTH (SQL):

MOLECULE TYPE (CI): DIVISION CODE (CI):

ORGANISM (ORGN):

AUTHOR (AU):

TITLE (TI):

JOURNAL (SO): OTHER SOURCE (OS):

AUTHOR (AU):

TITLE (TI):

FEATURES (FEAT): Feature Key

JOURNAL (SO):

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rGrb14
/protein-id="AAC61478.1"
/db-xref="GI:3650500"
/translation="MTTSLQDGQSAAGRAGAQDS
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LHALSPASDATRRGAMDRRKAKDLEVQETPSIPN
PFPELCCSPLTSVLSAGLFPRSNS
RKKQVIKVYSEDETSRALEVPSDVTARDVCQLLI
LKNHYVDDNSWTLFEHLSHTGVER
TVEDHELLTEVLSHWVMEEDNKLYLRKNYAKYEF
FKNPMYFFPEHMVSFATEMNGDRS
LTQIPQVFLSSNTYPEIHGFLHAKEQGKKSWKKA
YFFLRRSGLYFSTKGTSKEPRHLQ
FFSEFSTSNVYMSLAGKKKHGAPTPYGFCFKPTK
AGGPRDLKMLCAEEDQSRMCWVTA
IRLLKYGMQLYQNYMHPSQARSACSSQSVSPMRS
VSENSLVAMDFSGQKTRVIDNPTE
ALSVAVEEGLAWRKKGCLRLGNHGSPTAPSQSSA
VNMALHRSQPWFHHRISRDEAQQL
ITRQGPVDGVFLVRDSQSNPRTFVLSMSHGQKIK
HFQIIPVEDDGEVFHTLDDGHTKF
TDLIQLVEFYQLNKGVLPCKLKHYCARMAV"
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SEQUENCE (SEQ): 1 gctggacccc agcctttctt cgctttcgcc tcgcggtcga tgactcccta gaccccctgg 61 cctacgatca tgaccacgtc cctgcaagat gggcagagcg ccgcgggccg ggcgggcgcc 121 caggactccc cgctggcagt gcaggtgtgc cgcgttgccc agggcaaggg agacgcccag 181 gaččeggete aggtececgg actgeaegeg etgtececgg ceteagatge gaccegeege 241 ggtgccatgg acaggagaaa agcgaaagat ctggaagttc aggaaacgcc ttccattcct 301 aaccccttcc ctgagctctg ctgttctcca cttacatcgg tgctgtcagc aggcctcttc 361 cccagatcaa attcaaggaa gaaacaggtg attaaagttt acagcgagga tgagaccagc 421 agagcgttag aggtgcccag tgacgtcaca gcccgtgatg tctgccagct gttgatcctg 481 aagaaccact atgtcgacga caatagctgg accctttttg agcacctgtc tcacacaggc 541 gtagaaagga cagtggagga ccatgagctg ctgactgaag tgctgtctca ttgggtgatg 601 gaagaagata ataagctgta tcttagaaag aattatgcca aatatgaatt ttttaagaac 661 ccaatgtatt tctttccaga gcacatggtg tcttttgcaa ctgaaatgaa cggtgacaga 721 tecettacae agatecegea ggtgttttta ageteaaaea catateetga aatecatgge 781 ttcctgcatg caaaggaaca ggggaagaag tcttggaaaa aagcttactt ttttctcaga 841 agatctggtt tatatttttc tactaaaggc acatccaagg aaccacggca cttgcagttt 901 ttcagtgaat tcagcactag taatgtttac atgtcactgg caggcaaaaa aaagcatgga 961 gcgccgactc cctatggatt ctgctttaag cctaccaaag caggagggcc ccgggacctg 1021 aaaatgctgt gtgcagaaga agaccaaagc aggatgtgct gggtgaccgc cattagattg 1081 ctcaagtatg gcatgcagct ctaccagaat tatatgcatc catcccaagc tagaagcgcc 1141 tgcagītete agagēgtāte acceatgaga agegtāteag agaatteeet agīagēaātg 1201 gacttetcag gteagaagae cagagtéata gácáaceceá cigaagecet tieggitgeé 1261 gttgaggaag gactcgcttg gaggaaaaaa ggatgtttac gcctggggaa tcatgggagt 1321 cccactgcgc cctctcagag ctctgctgtg aacatggctc tccaccggtc ccagccatgg 1381 tttcaccaca gaatttctag agatgaagct cagcagttga ttacccggca ggggcctgtg 1441 gatggagttt tcttggtacg ggatagtcag agtaacccca gaacttttgt actgtcaatg 1501 agtcacggac aaaagataaa acactttcaa attatacccg tggaagatga tggtgaggtg 1561 ttccacaccc tggatgatgg ccatacgaag ttcacagatc tcatccagct cgtggagttc 1621 taccagctca acaaggggt ccttccttgc aagctgaagc attactgtgc taggatggct 1681 gtttagccaa actgicigig actcgitaaa ciatggaaga iggaggaigc aaagaagaat 1801 tcacaaggct ggaaacaaat catggtgaaa agaagattca cctgtgggtt acaaaaaaat 1861 aggtcacgta ttgcaaatta gtgaagactt ggattcgtat tactctcgtt actttaaatt 1921 tattagttaa aattaaacct tattaaaaaa

L6 ANSWER 130 OF 135 GENBANK.RTM. COPYRIGHT 2004 on STN

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LOCUS (LOC):
GenBank ACC. NO. (GBN): AI094433
GenBank VERSION (VER): AI094433.1 GI:3433409
CAS REGISTRY NO. (RN): 392191-42-3
SEQUENCE LENGTH (SQL): 420
MOLECULE TYPE (CI): mRNA; linear
DIVISION CODE (CI): Expressed sequence tag
DATE (DATE): 10 Nov 1998
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DEFINITION (DEF): ou87b07.s1 Soares_NSF_F8_9w_OT_PA_P_S1 Homo sapiens cDNA clone IMAGE:1634773 3' similar to TR:Q14449 Q14449

GRB14 . ;, mRNA sequence.

SOURCE:

ORGANISM (ORGN):

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;

```
Hominidae; Homo
NUCLEIC ACID COUNT (NA): 115 a
                                        75 c
                                                  67 g
                                                           163 t
COMMENT:
      Contact: Robert Strausberg, Ph.D.
      Email: cgapbs-r@mail.nih.gov
       This clone is available royalty-free through LLNL; contact the
       IMAGE Consortium (info@image.llnl.gov) for further information.
       Insert Length: 796
                                 Std Error: 0.00
      Seq primer: -40m13 fwd. ET from Amersham
      High quality sequence stop: 277.
REFERENCE:
                              1 (bases 1 to 420)
    AUTHOR (AU):
TITLE (TI):
                              NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
                              National Cancer Institute, Cancer Genome Anatomy
Project (CGAP), Tumor Gene Index
    JOURNAL (SO):
                              Unpublished (1997)
FEATURES (FEAT):
                                                      Qualifier
   Feature Key
                       Location
/organism="Homo sapiens"
/db-xref="taxon:9606"
/clone="IMAGE:1634773"
                    1..420
source
                                                  /clone-lib="Soares-NSF-F8-9W-OT-PA
                                                  -P-S1"
                                                  /lab-host="DH10B"
                                                  /note="Organ: pooled; Vector:
                                                  pT7T3D-Pač (Pharmacia) with a
                                                  modified polylinker; Site-1: Not
                                                  I; Site-2: Eco RI; Equal amounts
                                                  of plasmid DNA from five
                                                  normalized libraries were mixed,
                                                  and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a
                                                  subtractive hybridization
                                                  reaction. The driver was
                                                  PCR-amplified cDNAs from pools of
                                                  5,000 clones made from the same 5
                                                  libraries. The pools consisted of
                                                  the following libraries and
                                                  cloneIDs: Soares NbHSF pool 1: 309384-310919, 323208-325895
                                                  Soares Nb2HP pool 1:
145032-147335, 147720-148103,
148872-149255, 15002 - 150407,
151176-152327 Soares Nb2HF8-9W
                                                  pool 1: 758280-760583
                                                  772104-774407 Soares NbHPA pool 1:
                                                  304776-306311, 320136-322823, 326280-326663 Soares NbHOT pool 1:
                                                   723720-726407, 739080-740999
                                                  Subtraction by Bento Soares and M. Fatima Bonaldo."
SEQUENCE (SEQ):
      1 cctaaggttt aattttaact aatgaatttt aaatgatgaa tgtaaagtca atccaagtct
     61 ttgcttattt gcaatgcaca aactattttt ttgtäacttg cäggtgäaat acattctttt
    121 cacatgataa cgttttcgcc cttatttatg gtcttttatt attittcttg agtccttttc
181 cttcaatagt ttaataagtc acttctggct tgtctagaga gcaatcctag cacaataatg
    241 tttcaacttg caaggaagaa cgcccttatt gagttgatag aactccacca gctgtattag
301 atctgtaaat cttgtgtggc catcatccag tgtgtggaac atttcaccgt catcttctac
361 tggtataatt tgaaagtgct ttattttttg tccatgactc attgacagta cgaaagtttt
L6
      ANSWER 131 OF 135
                                GENBANK.RTM. COPYRIGHT 2004 on STN
LOCUS (LOC):
                              AA917917
                                              GenBank (R)
GenBank ACC. NO. (GBN): AA917917
GenBank VERSION (VER):
                              AA917917.1
                                            GI:3057807
CAS REGISTRY NO. (RN):
SEQUENCE LENGTH (SQL):
MOLECULE TYPE (CI):
                              206814-02-0
                              497
                              mRNA; linear
DIVISION CODE (CI):
                              Expressed sequence tag
DATE (DATE):
                              10 Jun 1998
DEFINITION (DEF):
                              ol76g09.s1 NCI_CGAP_Kid3 Homo sapiens cDNA clone
                              IMAGE: 1535584 3' similar to TR: Q14449 Q14449
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GRB14

. ;, mRNA sequence.

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SOURCE:
                            human.
 ORGANISM (ORGN):
                            Homo sapiens
                            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                            Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                            Hominidae; Homo
                                      95 c
                                             79 g
NUCLEIC ACID COUNT (NA): 136 a
                                                       187 t
COMMENT:
      Contact: Robert Strausberg, Ph.D.
      Email: cgapbs-r@mail.nih.gov
      Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
      Emmert-Buck, M.D., Ph.D.
       cDNA Library Preparation: M. Bento Soares, Ph.D.
       cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
       Clone distribution: NCI-CGAP clone distribution information can be
      found through the I.M.A.G.E. Consortium/LLNL at:
      www-bio.11n1.gov/bbrp/image/image.html
      Insert Length: 664
                              Std Error: 0.00
      Seq primer: -40m13 fwd. ET from Amersham
      High quality sequence stop: 440.
                            1 (bases 1 to 497)
REFERENCE:
   AUTHOR (AU):
                            NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
                            National Cancer Institute, Cancer Genome Anatomy
Project (CGAP), Tumor Gene Index
   TITLE (TI):
                            Unpublished (1997)
   JOURNAL (SO):
FEATURES (FEAT):
                                                   Qualifier
                     Location
  Feature Key
/organism="Homo sapiens"
/db-xref="taxon:9606"
/clone="IMAGE:1535584"
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source
                                               /lab-host="DH10B"
                                               /note="Organ: kidney; Vector:
                                               pT7T3D-Pac (Pharmacia) with a
                                               modified polylinker; Site-1: Not
                                               I; Site-2: Eco RI; 1st strand cDNA
                                               was primed with a Not I -
                                               oligo(dT) primer, double-stranded cDNA was ligated to Eco RI
                                              adaptors (Pharmacia), digested with Not I and cloned into the Not
                                               I and Eco RI sites of the
                                               modified pT7T3 vector. mRNA
                                               source: 2 pooled kidneys. Library
                                               went through one round of
                                              normalization. Library constructed by Bento Soares and M. Fatima
                                               Bonaldo.
SEQUENCE (SEQ):
      1 cattttttc ctaaggttta attttaacta atgaatttta aatgatgaat gtaaagtcaa
     61 tccaagtctt tgcttatttg caatgcacaa actattttt tgtaacttgc aggtgaaata
   121 cattettte acatgataat gttttegeee ttatttatgg tettttatta tittettga 181 gteetttee tteaatagtt taataagtea ettetggett gtetagagag caateetage
   241 acaataatgt ttcaacttgc aaggaagaac gcccttattg agttgataga actccaccag 301 ctgtattaga tctgtaaatc ttgtgtggcc atcatccagt gtgtggaaca tttcaccgtc 361 atcttctact ggtataattt gaaagtgctt tattttctgt ccatgactca ttgacagtac
   421 gaaagttttg gggttactct gactatcccg taccaagaaa actcatccac aagtccttgc
   481 tgaataatca atcgctg
L6
     ANSWER 132 OF 135
                              GENBANK.RTM. COPYRIGHT 2004 on STN
LOCUS (LOC):
                            AA684351
                                           GenBank (R)
GenBank ACC. NO. (GBN): AA684351
GenBank VERSION (VER):
                            AA684351.1 GI:2670937
CAS REGISTRY NO. (RN):
                            200792-61-6
SEQUENCE LENGTH (SQL):
                            503
MOLECULE TYPE (CI):
                            mRNA; linear
DIVISION CODE (CI):
                            Expressed sequence tag
DATE (DATE):
                            9 Dec 1997
DEFINITION (DEF):
                            vm69d04.s1 Knowles Solter mouse 2 cell Mus musculus
                            CDNA clone IMAGE:1003495 5' similar to TR:Q14449 Q14449
                              ***GRB14*** . ;, mRNA sequence.
SOURCE:
```

house mouse.

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ORGANISM (ORGN):
                            Mus musculus
                             Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                            Euteleostomi; Mammalia; Eutheria; Rodentia;
                            Sciurognathi; Muridae; Murinae; Mus
NUCLEIC ACID COUNT (NA): 127 a
                                                 135 g
                                       121 c
COMMENT:
      Contact: Marra M/Mouse EST Project
      WashU-HHMI Mouse EST Project
      Washington University School of MedicineP
      4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
      Tel: 314 286 1800
      Fax: 314 286 1810
      Email: mouseest@watson.wustl.edu
      This clone is available royalty-free through LLNL; contact the
      IMAGE Consortium (info@image.linl.gov) for further information.
      MGI:567711
      Possible reversed clone: similarity on wrong strand
      High quality sequence stop: 459.
                                (bases 1 to 503)
REFERENCE:
   AUTHOR (AU):
                            Marra,M.; Hillier,L.; Allen,M.; Bowles,M.; Dietrich,N.;
                            Dubuque,T.; Geisel,S.; Kucaba,T.; Lacy,M.; Le,M.; Martin,J.; Morris,M.; Schellenberg,K.; Steptoe,M.; Tan,F.; Underwood,K.; Moore,B.; Theising,B.; Wylie,T.; Lennon,G.; Soares,B.; Wilson,R.; Waterston,R.
                             The WashU-HHMI Mouse EST Project
   TITLE (TI):
   JOURNAL (SO):
                            Unpublished (1996)
FEATURES (FEAT):
                                                    Qualifier
                       Location
  Feature Key
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source
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/db-xref="taxon:10090"
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                                                /clone-lib="Knowles Solter mouse 2
                                                /tissue-type="embryo"
                                                /dev-stage="2-ce11
                                                /lab-hosť="DH10B'
                                                /note="Organ: embryo; Vector:
                                                pBluescribe (modified); Site-1:
                                                MluI; Site-2: SalI; Cloned unidirectionally from mRNA
                                                prepared from 13,500 2-cell stage
                                                embryos. Primer: SalI(dT):
                                                5'-CGGTCGACCGTCGACCGTTTTTTTTTTTTT
                                                         cDNAs were cloned into the
                                                MluI/SalI sites of a modified
                                                pBluescribe vector using
                                                commercial linkers (NEB). Average insert size: 1.2 kb."
SEQUENCE (SEQ):
      1 ttccctagta gcaatggact tctcaggtga gaagagcaga gtcatagaca accccactga
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    121 ggggaatcač ggaagcccca gtgccccctc ccagagctct gctgtgaaca tggctctcca
    181 tcggtcccaa ccatggtttc accacagaat ttccagagat gaggctcagc ggctgatcat
   241 tcggcagggg cctgtggatg gagttttctt ggtacgggat agtcagagta accccagaac 301 ttttgtactg tcaatgagtc atggacaaaa gataaaacac tatcaaatta tacccgtaga 361 agatgatggt gagctgttcc atactctgga tgatggccat acgaagttca cagacctcat 421 ccagctggtg gagttctacc agctcaacag gggggtcctt ccttgcaagc tgaagcatta
    481 ctgtgctagg atggctgttt agc
L6
      ANSWER 133 OF 135
                                                COPYRIGHT 2004 on STN
                               GENBANK.RTM.
LOCUS (LOC):
                            HSU69276
                                            GenBank (R)
GenBank ACC. NO. (GBN): U69276
GenBank VERSION (VER): U69276
                            U69276.1 GI:1546834
                             181109-72-8
CAS REGISTRY NO. (RN):
SEQUENCE LENGTH (SQL):
                            2504
MOLECULE TYPE (CI):
                            mRNA; linear
DIVISION CODE (CI):
                             Primates
DATE (DATE):
                             17 Sep 1996
DEFINITION (DEF):
                            Human hGrbIRbeta/hGrb10 (GRBIRbeta/GRB10) mRNA,
                             complete cds.
SOURCE:
                             human.
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                            Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                           Hominidae; Homo
                                    652 c
NUCLEIC ACID COUNT (NA): 639 a
                                               654 g
                                                         555 t 4 others
                           1 (bases 1 to 2504)
REFERENCE:
   AUTHOR (AU):
                           Frantz, J.D.; Giorgetti-Peraldi, S.; Ottinger, E.A.;
                           Shoelson, S.E.
   TITLE (TI):
                           Human GrbIRbeta/Grb10: Splice Variants of an Insulin
                           and Growth Factor Receptor-Binding Protein with PH and
                           SH2 Domains
                           Unpublished
   JOURNAL (SO):
                           2 (bases 1 to 2504)
Frantz, J.D.; Giorgetti-Peraldi, S.; Ottinger, E.A.;
REFERENCE:
   AUTHOR (AU):
                           Shoelson, S.E.
   TITLE (TI):
                           Direct Submission
                           Submitted (04-SEP-1996) Research Division, Joslin
   JOURNAL (SO):
                           Diabetes Center, One Joslin Place, Boston, MA 02215,
                           USA
FEATURES (FEAT):
  Feature Key Location
                                                   Qualifier
/organism="Homo sapiens"
/db-xref="taxon:9606"
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source
                                              /tissue_type="cerebellum and
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     AUTHOR (AU):
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                                          novel member of the GRB7 gene family J. Biol. Chem., 271 (21), 12502-12510 (1996)
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Dep. Cell Biol., Scripps Res. Inst., La Jolla, CA 92037, USA
American Society for Cell Biology, 9650 Rockville Pike, Bethesda, MD
20814, USA; phone: (301) 530-7153; fax: (301) 530-7139; email:
ascbinfo@ascb.org; URL: www.ascb.org/ascb/, Abstracts available. Price
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       Meeting (9840478). San Francisco, CA (USA). 12-16 Dec 1998. ASCB. Bio-Rad,
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